

KM treatment; *Neisseria* infection; meningitis; septicaemia; gonorrhea.
 XX
 OS *Neisseria meningitidis*.
 XX
 PN W09924578-A2.
 XX
 PD 20-MAY-1999.
 XX
 PF 09-OCT-1998; 98MO-IB01665.
 XX
 PR 01-SEP-1998; 98GB-0019016.
 PR 06-NOV-1997; 97GB-0023516.
 PR 14-NOV-1997; 97GB-0024190.
 PR 18-NOV-1997; 97GB-0024386.
 PR 27-NOV-1997; 97GB-0025158.
 PR 10-DEC-1997; 97GB-0026147.
 PR 14-JAN-1998; 98GB-0000759.
 XX
 PA (CHIR-) CHIRON SPA.
 XX
 PI Grandi G, Masignani V, Pizzo M, Rappuoli R, Scarlato V;
 XX WPI: 1999-327407/27.
 DR N-PSDB; AA212026.
 XX
 PT Proteins from *Neisseria meningitidis* and *N. gonorrhoeae* useful for
 PT diagnosis, treatment and prevention of infection
 XX
 PS Claim 4; Page 123; 524pp; English.
 XX
 CC Amino acid sequences AAY38499-Y38944 represent *Neisseria meningitidis*
 CC and *N. gonorrhoeae* antigenic proteins. They are encoded by open
 CC reading frames (ORFs) AA21972-212358. The antigenic proteins,
 CC their fragments, their nucleic acids and antibodies are used for
 CC diagnosis, prevention (as vaccines) or treatment of *Neisseria*
 CC infections, such as meningitis, septicaemia and gonorrhea. Both
 CC organisms are closely related. Fragments of the nucleic acids
 CC are useful as hybridisation probes and antisense reagents.
 XX
 SQ Sequence 447 AA;

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 Ratio: 5.121 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

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 201 GTTACTGCGCGGCTTCAGGCAAAATCGCGGATTCACCGTGGCGAAA 250
 67 lphenhrilaproalaserglyllysllealalalelhisarglyllyl 84
 251 AGCGGCTACTGACTGATCGTGTGCGTGAAGGCAACGAGGAATC 300

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 101 GluphegluarglylralprogluvalaleuallaasnleuSerGlyllyl 117
 351 AGTGGCGCGCAACCTGATCCAAATCCGGTGTGAGCTGGCGTGCACCC 400
 117 uvalalargatgasnleuileglinserclyleuphrilaleuarglylra 134
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 134 rpropheserlyslleproalalvalaspalagluprophealalephe 150
 451 GTCATGCGATGACACCAATCCGCGTGGCGGCGGCGGCGGATCAT 500
 151 ValasnalemetaspthrlnsProleuallaalaspProthrallel 167
 501 CAAGAAGCGCGCGAGATTCAAACGCGCGCTGTGTGATTAAGCCGTT 550
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seq_documentation_block:
ID   AAV38562 standard; Protein; 447 AA.
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AC   AAV38562;
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DT   08-OCT-1999 (first entry)
XX
DE   Neisseria meningitidis strain A antigen encoded by ORF22.
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KW   Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
      treatment; Neisseria infection; meningitis; septicemia; gonorrhea.
XX
OS   Neisseria meningitidis.
XX
PN   WO924578-A2.
XX
PD   20-MAY-1999.
XX
PF   09-OCT-1998; 98WO-IB01665.
XX
PR   01-SEP-1998; 98GB-0019016.
      06-NOV-1997; 97GB-0023516.
      14-NOV-1997; 97GB-0024190.
      18-NOV-1997; 97GB-0024386.
      27-NOV-1997; 97GB-0025158.
      10-DEC-1997; 97GB-0026147.
      14-JAN-1998; 98GB-0000759.
XX
PA   (CHIR-) CHIRON SPA.
XX
PI   Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;

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XX
DR   WPI: 1999-327407/27.
DR   N-PSDB; AA12027.
XX
PT   Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
      diagnosis, treatment and prevention of infection
XX
PS   Claim 4; Page 123; 524pp; English.
XX
CC   Amino acid sequences AAV38499-Y38944 represent Neisseria meningitidis
      and N. gonorrhoeae antigenic proteins. They are encoded by open
      reading frames (ORFs) AA11972-Z12358. The antigenic proteins,
      their fragments, their nucleic acids and antibodies are used for
      diagnosis, prevention (as vaccines) and treatment of Neisseria
      infections, such as meningitis, septicemia and gonorrhea. Both
      organisms are closely related. Fragments of the nucleic acids
      are useful as hybridisation probes and antisense reagents.
XX
SQ   Sequence 447 AA:

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      Ratio: 4.982          Gaps: 0
      Percent Similarity: 97.763      Percent Identity: 94.855

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51 GCAAGCGGTTTACGACGCGCGCGCATTAACGGAAGTGGCGTTGGCG 100
  17 uGlnValIleTyrAspGlyProValIleThrGluValAlaLeuLeuGly 34
101 AAGATATATGCGGTATGCGCGCCCTCGATGAAGTCAAGGAAGCCATGC 150
  34 lngIulYrAlaGlyMetArgPro**MetLysValLysGluLysAla 50
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  51 ValLysLysGlyGlnValleuPheGluAspLysLys**ProGlyValVa 67
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  117 u*****AsnLeuIleGlnSerGlyLeuTrpThrAlaLeuArg***A 134
401 GTCGTTTCAGCAAAATTCCTGCGCGATGCGGAGCGCGGTGCGCATPCT 450
  134 rGProPheSerLysIleProAlaValaAspAlaGluProPheAlaIlePhe 150
451 GTCAATGCGATGACACCAATCCGCTGCTGCGGACCTACCGCATATAT 500
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501 CAAAGAGCGCGCGAGGATTTCAAAACGGCGGCTGTGATTGATGACCGGT 550
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601 TCTGAAATGCTGCCAATCGAATCGAAGACATGAAATTCGGCGCGCATCC 650
201 SerGluSnAlaAlaSnIleGluThrHisGluPheGlyGlyProHisSpr 217
651 TGCCGGTTTGAGTGGCAGCAGCATTCATTCATGACCGCGTGGCGCGA 700
217 oAlaGlyLeuSerGlyThrHisIleHisPheIleGluProValIleVala 234
701 ATAAACCGGTGGACCATCAATATCAAGATGTAATTCACCATGGCGGT 750
234 snlysrhValTrpThrIleAsnTrpGlnAspValIleAlaIleGlyArg 250
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251 LeuPheAlaThrGlyArgLeuAsnThrGluArgValIleAlaIleGly 267
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401 GlyAspThrAspSerAlaGlnAlaIleuGlyCysLeuGluLeuAspGlu 417
1251 AGACCTGCTTTGTCAGCTTCTGTCGCGCGCGCAATACGATACGCC 1300
417 uAspLeuAlaIleuCysSerPheValCysProGlyLysTrpGlu**Gly 434
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seq_documentation_block:

ID AAY38563 standard; protein; 322 AA.

XX AAY38563;

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DT 08-OCT-1999 (first entry)

DE Neisseria gonorrhoeae antigen encoded by a partial ORF22.

XX Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;

KW treatment; Neisseria infection; meningitis; septicemia; gonorrhea.

OS Neisseria gonorrhoeae.

XX W09924578-A2.

XX 20-MAY-1999.

XX 09-OCT-1998; 98W0-IB01665.

XX 01-SEP-1998; 98GB-0019016.

XX 06-NOV-1997; 97GB-0023516.

XX 14-NOV-1997; 97GB-0024190.

XX 18-NOV-1997; 97GB-0024386.

XX 27-NOV-1997; 97GB-0025158.

XX 10-DEC-1997; 97GB-0026147.

XX 14-JAN-1998; 98GB-0000759.

XX (CHIR-) CHIRON SPA.

XX Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;

PI WPI: 1999-327407/27.

XX proteins from Neisseria meningitidis and N. gonorrhoeae useful for

PT diagnosis, treatment and prevention of infection

XX Claim 4; Page 124-125; 524pp; English.

XX Amino acid sequences AAY38499-Y38944 represent Neisseria meningitidis

CC and N. gonorrhoeae antigenic proteins. They are encoded by open

CC reading frames (ORFs) AA211972-212358. The antigenic proteins,

CC their fragments, their nucleic acids and antibodies are used for

CC diagnosis, prevention (as vaccines) or treatment of Neisseria

CC infections, such as meningitis, septicemia and gonorrhea. Both

CC organisms are closely related. Fragments of the nucleic acids

CC are useful as hybridisation probes and antisense reagents.

XX Sequence 322 AA;

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Quality: 1576.00 Length: 322

Ratio: 4.910 Gaps: 0

Percent Similarity: 99.689 Percent Identity: 95.031

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17 uGlnValIleTrpAspGlyProAlaIleThrGluValAlaLeuLeuGly 34

101 AAGCATATGCCGGTATCGCCCTCGCATGAAAGTCAAGAGAGCGATGCC 150

34 IuGluTrpValGlyMetArgProSerMetLysIleLysGluIuGlyAla 50

151 GTCAAAAAGCCCAAGTCTGTTTGAAGCAAAAAGAAATCCGGCGTGT 200

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401 GTCGCGTTCGCAAAATTCCTGGCGTCGATCCGACGCGCGTGGCGATGTC 450
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seq_documentation_block:
ID: AA198560 standard; Protein; 158 AA.
XX AA198560;
XX AC
XX DT
XX 08-Oct-1999 (first entry)
Neisseria meningitidis antigen encoded by a partial ORF22.

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seq_documentation_block:
ID   AAV34439 standard; Protein; 451 AA.
XX
AC   AAV34439;
XX
DT   25-AUG-1999 (first entry)
XX
DE   Porphyromonas gingivalis protein PGI.
XX
KM   Porphyromonas gingivalis; Pg; periodontal disease; gingivitis;
XX   vaccine; antigenic.
XX
OS   Porphyromonas gingivalis.
XX
WO929870-A1.
XX
PN   17-JUN-1999.
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PD   10-DEC-1998; 98WO-AU01023.
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PF   04-AUG-1998; 98AU-0005028.
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XX   31-DEC-1997; 97AU-0001182.
XX   30-JAN-1998; 98AU-0001546.
XX   10-MAR-1998; 98AU-0002264.
XX   09-APR-1998; 98AU-0002911.
XX   23-APR-1998; 98AU-0003128.
XX   05-MAY-1998; 98AU-0003338.
XX   22-JUL-1998; 98AU-0003654.
XX   29-JUL-1998; 98AU-0004917.
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XX   Ross BC, Rothel LJ, Webb EA;
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DR   WPI, 1999-385613/32.
XX   N-PSDB; AAV91657.
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PT   Antigenic Porphyromonas gingivalis peptides for preventing
XX   gingivitis
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PS   Claim 1; Page 417-418; 588pp; English.
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CC   AAX91536 to AAX91801 encode two hundred and sixty six antigenic
CC   Porphyromonas gingivalis (Pg) polypeptide sequences given in AAV34318 to
CC   AAV34583. AAX91802 to AAX91989 represent PCR primers used in the
CC   isolation of the Pg polypeptides. The Pg polypeptides have antibacterial
CC   activity with a vaccine mechanism of action. The Pg polypeptides can be
CC   used as vaccines especially against Porphyromonas gingivalis. Probes can

```

CC be used to detect Porphyromonas gingivalis in standard hybridisation
 CC assays. Porphyromonas gingivalis is involved in periodontal disease
 CC especially gingivitis.

XX SQ Sequence 451 AA:

alignment_scores:
 Quality: 663.00 Length: 452
 Ratio: 2.225 Gaps: 7
 Percent Similarity: 65.929 Percent Identity: 34.735

alignment_block:

US-09-303-518D-125 x AAV34439

Align seg 1/1 to: AAV34439 from: 1 to: 451

```

1 ATGATTAAATCAAAAAAGTCTAAACCTGCCATCGCGGACGACGGA 50
  ::::: ||||| ||||||| ||| :::: ||::: |||
4 ValIleLysThrLysLysGlyLeuAlaLeuAsnLeuLysGlyProIle 20
  ::::: ||||| ||||| ::::: |||
51 GCAGCGCGTTTACGACGCGCGCCATTTACCGAAGTC...GCGTGGCTTG 97
  ::::: ||||| ::::: |||
20 uProGluMetLeuAlaGluProAlaGlnSerProThrTyrAlaValAlp 37
  ::::: ||||| ||||| ::::: |||
98 GCGAGATATATGCCGCTATGCCGCCCTCGATGAAAGTCAAGGACGAT 147
  ::::: ||||| ||||| ::::: |||
37 roAspAspPheGluGlyValIleProLysValThrAlaTrpGluLysp 53
  ::::: ||||| ||||| ::::: |||
148 GCCGTCAAAAAAGGCCCAAGTGTGTTTGAAGACAAAAAGATCCGGCGT 197
  ||::: ||::: ||::: ||::: ||::: |||
54 LysValAlaGlaGlySerAlaLeuMetHisLysAlaTyrProGluMet 70
  ::::: ||||| ||||| ::::: |||
198 GGTGTTTACTGCGCGCGCTTCAAGCAAAATCCCGGATTCACCGTGCG 247
  ::::: ||||| ||||| ::::: |||
70 LysPheThrSerProValSerGlyGluValIleAlaValAsnArgGlyA 87
  ::::: ||||| ||||| ::::: |||
248 AAAGCGCGTACTCAGTCAGTCGATGCGCGCTTGAAGCAGCAGCA 297
  ||||| ::::: ||||| ::::: |||
87 LysArGlyLysValLeuSerIleGluValLysProkspGlyLeuAsnGlu 103
  ::::: ||||| ||||| ::::: |||
298 ATCGAG...TTTGAAAGCTACGACCTGAGCGCTGGCAACTTAAGCG 344
  ||| ||| ||||| ||||| |||||
104 TyrGluSerPheProValGlyLysProSerAla.....LeuSerAl 117
  ::::: ||||| ||||| ::::: |||
345 CGAAGAGTGGCGCGCGACCTGATCCATCCGCTTGTGACTGCGGTGC 394
  ::::: ||||| ||||| ::::: |||
117 agLInIleLysGluLeuLeuSerSerGlyMetTrpGlyPheIle 134
  ::::: ||||| ||||| ::::: |||
395 GCACCGCGCTTCAGCAAAATTCCTGCGCGTGGATGCCGACCGCTTCC 444
  ::::: ||||| ||||| ::::: |||
134 yseGlnArProIlyTrsPrlLeuAlaIleThrProAspIleAlaProArgasp 150
  ::::: ||||| ||||| ::::: |||
445 ATCTTGCAATGCGATGACGACCAATCCGCTGCTGCGGACCTTAACG 494
  ||::: ||::: ||::: ||::: ||::: |||
151 IleTyrIleThrAlaAsnPheThrAlaProLeuAlaIleProAspPheAsp 167
  ::::: ||||| ||||| ::::: |||
495 CATTTCAAGAAAGCGCGCGGAGATTTCAAACGCGCGCTGTTGATTGA 544
  ::::: ||||| ||||| ::::: |||
167 eIleValArgGlyGluGluArgAlaLeuGlnThrAlaIleAspAlaLeu 184
  ::::: ||||| ||||| ::::: |||
545 GCGGTTTGACGACGACGCAAAATTCATGTTTGAAGCAGCTGCGGCAAC 594
  ::::: ||||| ||||| ::::: |||
184 LysLeuThrThrLysValTyrValGlyLeuLysProLysSer 200
  ::::: ||||| ||||| ::::: |||
595 GTCCGCTGAAATGCTGCCAATCGAACACATGATTCGCGCGCC 644
  ::::: ||||| ||||| ::::: |||
201 LeuGlyLeuHisAsnAlaGluIleValGluHis.....GlyPr 214
  ::::: ||||| ||||| ::::: |||
645 GCATCTGCGCGTTGAGTGGACGACCAATTCATTCACGACGCGTGC 694
  ::::: ||||| ||||| ::::: |||
214 oHisProAlaGlyAsnValGlyValLeuIleAsnHisThrLysProIleA 231

```


CC Neisserial bacteria (e.g. meningitis and septicemia), to detect the
 CC presence of Neisseria bacteria, or to raise antibodies. They may also
 CC be used to screen for agonists or antagonists, which may themselves
 CC have use as antibacterial agents. The polynucleotides of the invention
 CC may also be used in gene therapy protocols.

XX Sequence 120 AA;

alignment_scores:

Quality: 600.00 Length: 120
 Ratio: 5.000 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-303-518D-125/rev x AAY75272

Align seg 1/1 to: AAY75272 from: 1 to: 120

```

674 ATGTGCGTCCACTCAACCGGAGATGGGGCCGCCGAATTCATGTGT 625
1 MetCysValProLeuLysProAlaGlyCysGlyProPheSerCysVal 17
624 TTGCATGTTGGCAGATTTCAGACGGCAGCTCTCCGACCTGCTTAC 575
17 IserMetLeuAlaAlaPheSerAspGlyThrSerAlaProAlaLeuG 34
574 AAACATGATTTTGGCTTGGTCAACCGCTCAATACCAACAGCCGGCGT 525
34 IntThrPheLeuAlaArgSerValLysArgLeuAsnThrSerAlaProArg 50
524 TTGAATCCTCGGGCGCTCTTTGATAATGACCGTAGGGTCGACGACG 475
51 LeuYssSerSerAlaAlaSerLeuIleMetThrValGlySerAlaAla 67
474 CGGATGTTGTCATCGCATTCAGACGATGCGCAAGCGCTCGGACATCG 425
67 rGlyLeuValSerIleAlaLeuThrLysMetAlaAsnGlySerAlaSer 84
424 CGGAGAGATTTTGTCTGAACGAGCGGGTGCAGCGCAGTCACAAACG 375
84 hrAlaGlyIleLeuLeuAlaAsnGlyArgValAlaArgSerAlaValHisLysPro 100
374 GATTGGATCAGGTTGGCGGCGACTTCTTCCGCCCTTAAGTTGGCAGCG 325
101 AspThrPheLeuAlaArgArgThrSerSerProLeuLysPheAlaSerAl 117
324 TTCAGGTGCG 315
117 aserGlyAla 120

```

seq_name: /SIDSI/gcgdata/geneseq/geneseqp-emb1/AA2000.DAT: AAY75273

seq_documentation_block:

ID AAY75273 standard: protein: 120 AA.

XX AAY75273;
 XX
 XX 21-MAR-2000 (first entry)
 XX
 DE Neisseria meningitidis ORF 628 protein sequence SEQ ID NO:2020.
 KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
 KW antigenic; diagnosis; immunogenic; infection; meningitis; septicemia;
 KW antibacterial; gene therapy.
 XX
 OS Neisseria meningitidis.
 XX
 PN WO957280-A2.
 XX
 PD 11-NOV-1999.
 XX
 PF 30-APR-1999; 99WO-US09346.

XX
 PR 01-MAY-1998; 98US-0083758.
 PR 31-JUL-1998; 98US-0094869.
 PR 02-SEP-1998; 98US-0098994.
 PR 02-SEP-1998; 98US-0098994.
 PR 09-OCT-1998; 98US-0103749.
 PR 09-OCT-1998; 98US-0103749.
 PR 09-OCT-1998; 98US-0103749.
 PR 25-FEB-1999; 99US-0121520.
 XX
 PA (CHIR) CHIRON CORP.
 PA (GENO-) INST GENOMIC RES.

PI Fraser C, Galeotti C, Grandi G, Hickey E, Masignani V, Mora M,
 PI Petersen J, Pizzo M, Rappuoli R, Ratti G, Scalato E, Scarselli M,
 PI Tettelin H, Venter JC;
 XX
 DR WPI; 2000-062150/05.
 DR N-PSDB; AA254035.

PT Novel Neisserial polypeptides predicted to be useful antigens for
 PT vaccines and diagnostics
 PS Claim 2; Page 1004; 1453pp; English.

CC AA253015 to AA254536, AA254577 to AA254615, and AAY74253 to AAY75941
 CC represent novel Neisseria meningitidis and N. gonorrhoeae polynucleotides
 CC and polypeptides. AA255537 to AA254576 and AA254616 to AA255473 represent
 CC PCR primers used in the exemplification of the present invention. The
 CC polypeptides, the polynucleotides, antibodies and compositions of
 CC the invention can be used as vaccines, as diagnostic reagents, and as
 CC immunogenic compositions. The polypeptides can be used in the
 CC manufacture of medicaments for treating or preventing infection due to
 CC Neisserial bacteria (e.g. meningitis and septicemia), to detect the
 CC presence of Neisseria bacteria, or to raise antibodies. They may also
 CC be used to screen for agonists or antagonists, which may themselves
 CC have use as antibacterial agents. The polynucleotides of the invention
 CC may also be used in gene therapy protocols.

XX Sequence 120 AA;

alignment_scores:
 Quality: 574.00 Length: 120
 Ratio: 4.824 Gaps: 0
 Percent Similarity: 99.167 Percent Identity: 95.000

alignment_block:

US-09-303-518D-125/rev x AAY75273

Align seg 1/1 to: AAY75273 from: 1 to: 120

```

674 ATGTGCGTCCACTCAACCGGAGATGGGGCCGCCGAATTCATGTGT 625
1 MetCysValProLeuLysProAlaGlyCysGlyProPheSerCysVal 17
624 TTGCATGTTGGCAGATTTCAGACGGCAGCTCTCCGACCTGCTTAC 575
17 IserMetLeuAlaAlaPheSerAspGlyThrSerAlaProAlaLeuH 34
574 AAACATGATTTTGGCTTGGTCAACCGCTCAATACCAACAGCCGGCGT 525
34 IntThrPheLeuAlaArgSerValLysArgLeuAsnThrSerLysProArg 50
524 TTGAATCCTCGGGCGCTCTTTGATAATGACCGTAGGGTCGACGACG 475
51 LeuYssSerSerAlaAlaSerLeuIleMetThrValGlySerAlaAla 67
474 CGGATGTTGTCATCGCATTCAGACGATGCGCAAGCGCTCGGACATCG 425
67 rGlyLeuValSerIleAlaLeuThrLysMetAlaAsnGlySerAlaSer 84
424 CGGAGAGATTTTGTCTGAACGAGCGGGTGCAGCGCAGTCACAAACG 375

```

```

|||||
84 hrhlaaglylleuleuansnilyarvalargseralavahlslypro 100
374 GATTGGATCAGGTTGGCGGCACCTTCTTCGCCGCTTAAGTTGGCAGCGC 325
|||||
101 AsprtlleargleuArgThrSerSerProleuLysPheAlaAsnAl 117
324 TTCAGGTGCG 315
|||||
117 aserGlyAla 120

seq_name: /SIDS1/gcgdata/geneseq/geneseqp-emb1/AA2000.DAT:AAV75271
seq_documentation_block:
ID AAV75271 standard; Protein; 119 AA.
XX
AC AAV75271;
XX
DT 21-MAR-2000 (first entry)
XX
DE Neisseria gonorrhoeae ORF 628 protein sequence SEQ ID NO:2016.
XX
KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
KW antigenic; diagnosis; immunogenic; infection; meningitis; septicemia;
KW antibacterial; gene therapy.
XX
OS Neisseria gonorrhoeae.
XX
PN WO957280-A2.
PD 11-NOV-1999.
XX
PF 30-APR-1999; 99WO-US09346.
XX
PR 01-MAY-1998; 98US-0083758.
PR 31-JUL-1998; 98US-0094869.
PR 02-SEP-1998; 98US-0098994.
PR 02-SEP-1998; 98US-0099062.
PR 09-OCT-1998; 98US-0103749.
PR 09-OCT-1998; 98US-0103794.
PR 09-OCT-1998; 98US-0103796.
PR 25-FEB-1999; 99US-0121528.
XX
PA (CHIR ) CHIRON CORP.
PA (GENO-) INST GENOMIC RES.
XX
PI Fraser C, Galeotti C, Grandi G, Hickey E, Maignani V, Mora M;
PI Petersen J, Piza M, Rappuoli R, Ratti G, Scalato E, Scarselli M;
PI Tettelin H, Venter JC;
XX
DR WPI: 2000-062150/05.
DR N-PSDB; AA254033.
XX
PT Novel Neisserial polypeptides predicted to be useful antigens for
PT vaccines and diagnostics
XX
PS Claim 2; Page 1003; 1453pp; English.
XX
AA253015 to AA254536, AA254577 to AA254615, and AAV74253 to AAV75941
CC represent novel Neisseria meningitidis and N. gonorrhoeae polynucleotides
CC and polypeptides. AA254537 to AA254576 and AA254616 to AA25473 represent
CC PCR primers used in the exemplification of the present invention. The
CC polypeptides, the polynucleotides, antibodies and compositions of
CC the invention can be used as vaccines, as diagnostic reagents, and as
CC immunogen compositions. The polypeptides can be used in the
CC manufacture of medicaments for treating or preventing infection due to
CC Neisserial bacteria (e.g. meningitis and septicemia), to detect the
CC presence of Neisseria bacteria, or to raise antibodies. They may also
CC be used to screen for agonists or antagonists, which may themselves
CC have use as antibacterial agents. The polynucleotides of the invention
CC may also be used in gene therapy protocols.
XX
SQ Sequence 119 AA:

```

```

alignment_scores:
Quality: 538.50 Length: 119
Ratio: 4.642 Gaps: 1
Percent Similarity: 97.479 Percent Identity: 93.277

alignment_block:
US-09-303-518D-125/rev x AAV75271 ..

Align seg 1/1 to: AAV75271 from: 1 to: 119

674 ATGTGGTGGCCACTCAACCGCAGGATGGCGGCGCCGCAATTCATGTGT 625
|||||
1 MetCysValProleuLysProAlaGlyCysGlyProProAsnSerCysVal 17
624 TTCGATGTTGGCAGCATTTTTCAGAGCGCAGCTGTGCGCCAGCTTTCAC 575
|||||
17 lserlleuAlaAlaPheSerAspGlyThrSerAlaProAlaAlaLeuH 34
574 AATCATGATTTTTCGTTGCTGTCGTCGTCGTCGTCGTCGTCGTCGTCGTC 525
|||||
34 lstrtrpilleuValSerValAlaArgleuAsnThrSnaArgProAlaG 50
524 TTGAAATCTCTGGCGGCTTCTTGATATATGACCGTAGGTTGCGCAGCAG 475
|||||
51 leuLysSerSerAlaAlaSerleuMetMetThrValGlySerAlaAla 67
474 CGGATTTGGTTCATCGCATTCGACGAAGATGGCGGAACGGCTCGGCATCA 425
|||||
67 rglyleuValSerlleAlaLeuThrLysMetAlaAsnGlySerAlaSerT 84
424 CGGAGGAAATTTTTCGTCGAACGCGAGGCTGGCGCAGCGCAGCAACCG 375
|||||
84 hrhlaaglylleuleuansnilyarvalargseralavahlslypro 100
374 GATTGGATCAGGTTGGCGGCACCTTCTTCGCCGCTTAAGTTGGCAGCGC 325
|||||
101 Asp...lleArgleuArgThrPheSerleuLeuAsnPhelaAlaSerAl 116
324 TTCAGGT 318
|||||
116 aserGly 118

seq_name: /SIDS1/gcgdata/geneseq/geneseqp-emb1/AA2000.DAT:AAV82082
seq_documentation_block:
ID AAV82082 standard; Protein; 467 AA.
XX
AC AAV82082;
XX
DT 01-JUN-2000 (first entry)
XX
DE Chlamydia pneumoniae antigen CPN100605 protein SEQ ID NO:2.
XX
KW Chlamydia pneumoniae; antigen; CPN100605 protein; immunisation;
KW vaccine; infection; antibacterial; antiinflammatory; bronchitis;
KW community acquired pneumoniae; upper respiratory tract infection;
KW sinusitis.
XX
OS Chlamydia pneumoniae.
XX
PN WO200006742-A2.
PD 10-FEB-2000.
XX
PF 27-JUL-1999; 99WO-IB01331.
XX
PR 27-JUL-1998; 98US-0094195.
PR 26-JUN-1999; 99US-0361443.
XX
PA (CONN-) CONNAUGHT LAB LTD.
XX

```

PI Murdin AD, Oomen RP;
 XX WPI: 2000-205466/18.
 DR N-PSDB: AAZ95378.
 XX

PT Chlamydia pneumoniae antigens used for immunization and protection
 against Chlamydia diseases -
 XX

PS Claim 6; Fig 1; 48bp; English.

CC The present sequence represents the Chlamydia pneumoniae antigen
 CC CPN100605 protein. The CPN100605 protein has antibacterial and
 CC antiinflammatory activities. The Chlamydia pneumoniae CPN100605
 CC polynucleotide and protein can be used in vaccination methods for
 CC preventing and treating Chlamydia infection (e.g. infections caused by
 CC C. trachomatis, C. psittaci, C. pneumoniae or C. pecorum). The
 CC polynucleotide can be used to produce the protein recombinantly, in the
 CC construction of vaccine vectors, as a vaccine agent, and in the
 CC construction of an attenuated Chlamydia strain. The protein are also be
 CC useful as a vaccine agent, and for the preparation of medicaments for
 CC treating or preventing Chlamydia infection, e.g. community acquired
 CC pneumonia, and upper respiratory tract infections such as bronchitis and
 CC sinusitis.
 XX

SO Sequence 467 AA;

alignment_scores:

Quality: 464.50 Length: 464
 Ratio: 1.585 Gaps: 15
 Percent Similarity: 63.147 Percent Identity: 30.172

alignment_block:

US-09-303-518D-125 x AAY82082 ..

Align seg 1/1 to: AAY82082 from: 1 to: 467

```

4 ATTAATAATCAAAAAAGTCTAAACCTGCCATCGCGGACAGACCG...GA 50
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
3 IletlrValasnaarglyLeuaspLeuSerleuInglySerProlysgl 19
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
51 GCAAGCGGTTTACGAC.....GGCCCGCATACCGAAGTCCGTTGC 94
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
19 uSerlyPheTyrasnlyslleasprGluPheValSerlleasprleu. 35
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
95 TTGGCGAAGAAATATGCCGCTGACGCCCTCGATGAAGTCAAGAGAGC 144
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
36 .....ArgProPheGlnProleuSerleuLysleuLysValGlnGlnGly 50
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
145 GATGCCGTCAAAAAGGCCAAGTGTGTTTAAGACAAAAAAGAAATCCGGG 194
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
51 AspalavalCysSerGlyAlaProIlealagLutryLysHisPheProas 67
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
195 CGTGTGTTTACTGCGCGGCTTCAGGCAAAATGCGCGCATTCACCGTG 244
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
67 nThrTyrIleThrSerHisValSerGlyValAlaThrAlaIleArgArg 84
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
245 GCGAAACCGCGTACTCTGACCTGCTGAT...GCCGTGAAGGCAAC 291
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
84 LyaSnlySargSerleuAspValIlelleLysLysThrProGlyPro 100
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
292 GACGAATCGAGTTTGAAGCTGACGACCTGAACGCTGGCAACTTAAG 341
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
101 ThrSerThrGlnLyr.....ThrtyrAspLeuGlnThrLeuSe 113
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
342 CGGCGAAGAAATGCGCGCAACTGATCAATCGCTTGGGACGTCGCG 391
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
113 rArgSerAspLeuSerGlnIlePheLysGlnAsnGlyLeuPheAlaLeu 130
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
392 TGCGCACCGCTCGCTTGAAGAAATTCCTCGCGTGCATGCCGAG...CCG 438
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
130 IeLysGlnArgProPheasp...IleProAlaIleProThrGlnThrPro 145
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :

```

```

439 TTGCGCATCTTGTGCAATGCGATGACACCAATCCGCTGCGCGACCC 488
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
146 ArgAspValAlaPheIleAsnLeuAlaAspAsnArgProPheThrProSer 162
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
489 TACGGTCATTATC.....AAGAAGCCCGCCAGGATP 520
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
162 oGluLysHisLeuAlaLeuPheSerSerArgGlnGlnGlyPheTyrValP 179
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
521 TCAACCGCGCGCTGTGTATGACCGCTTGACCCGACCAAAATTCAT 570
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
179 heValValGlyValArgAlaIleAlaLysLeuPheGlyLeuArgProHis 195
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
571 GTTGTGAAGCAGCTGCGCGACAGCTGCGCTGAAGATCTGCAACAT 620
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
196 lleValPheArgAspArgLeuThrPheProThrGlnGlyLeuLysThrI 212
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
621 C...GAACACATGAAATTCGCGCGCGGACATCTGCGGTTTGACATGCA 667
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
212 eAlaHisLeuHisThrValSerGlyProPheProSerGlySerProSerI 229
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
668 CGACATTCATTTATTCGACCGCGCGCGCGGCAATTA...ACCGTGTG 714
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
229 leHisIleHisSerValAlaProIleThrAsnGlnLysGlnValAlaPhe 245
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
715 ACCATCAATATGACAGATGTAATGATGACATTCGCGCTTGTGCAACAG 764
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
246 ThrLeuSerPheGlnAspValLeuThrIleGlnLysLeuPheLeuLysG 262
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
765 CGCTGTGAACACGACGCGCGTGAATGCGCTAGCGTGTGTCATGCAACA 814
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
262 YArgIleLeuHisGlnGlnValThrAlaLeuAlaGlyThrAlaLeuLys 279
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
815 AACCG.....CGGCTGTGCGTACCGCTTTGGGTGCGCAAGTATGSCAA 858
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
279 eSerLeuArgArgTyrValIleThrThrLysGlyLysSerPheSer 295
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
859 ...ATTATCGCGCGCAATGTTGTTGACACAGACAAACGCGGTGATTTCCG 905
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
296 LeuIleAsnLeuAsnAspIleSerAspAsnArgThr...LeuIleSerG 311
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
906 TTGCGTATTTGAACGCGCGCATTTACACAGCGCGCGACGAT...TATTTGG 952
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
311 YAspProLeuThrGlnArgLeuLysLysGlnGlnGlnProPheLeuG 328
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
953 GACGTACCAACATGATTCGTTATGCAAGAGCGCGCGCAAGAG 1002
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
328 LyrPheArgAspHisSerIleSerValLeuHisAsnProThrLysArgL 344
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
1003 CTGTTCGCGCTGGGTTGCGCGGACGCGGCAAAATCTCCATGACGCGTAC 1052
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
345 LeuPheSerPheLeuArgIleGlyPheAsnLysProThrPheThrLys 361
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
1053 AACCTCGCGCATTTCTGAAAAACAAACTCTTCAAGTTCAAC.....A 1096
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
361 rTyrLeuSerGlyPhePheLysLysLysArgThrTyrThrAsnProAsp 378
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
1097 CAGCGCTCAACGCGCGCGACCGCGCATGTCGCCATGTGCTACTACGAG 1146
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
378 hAsnLeuHisGlnGlnThrArgProIleIleAspThrAspIleLysArg 394
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
1147 CGCGTATGCGCTGGATATCTGCGCACCGCGCTGTTGGCGGATTAAT 1196
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
395 LysValMetProMetArgIleProValValProleuIleLysAlaValI 411
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
1197 GGTGCGGATACGACGCGCGCGACGATGCGTGTGCTGGAATGAGCG 1246
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
411 eThrLysAsnPheAspLeuAlaAsnGlnLeuGlyPheLeuGlnValCysG 428
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
1247 AAGAAGACCTCGCTTGTGCAAGCTTGTGCGCGCGCAATGACATAC 1296
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
428 LysGlnAspPheAlaLeuProThrLeuIleAspProSerLysThrGlnMet 444
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :
1297 GCGCGCGCTGTTGCGCAAGTGTGCAAAACATTTGAGAAGAA 1338
  ||| : : : : : ||| : : : : : ||| : : : : : ||| : : : : :

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70 nllhlsrserValserGlyValleuLyslleaspaValTYRA 87
265 .....TCAGTCGATGTCGGTTGAGGCGAC 291
87 spserSerGlyTYrProLysProAlaValPheIleSerValleuLysasp 103
292 GACGAATCGAGCTTGAACGCTACGACCGTGAAGCGCTG.....GCAAA 335
104 GIUTPrGIUGLylleaspaSerProAlaValleuLysGluCysAs 120
336 CTTAGCGCGGAGAGAGTGGCGCGCAACCTGATCCATCCGCTTGTGGA 385
120 nleuAspAlaLysGluIleValAlaLyslleSerAlaIleGlyle...V 136
386 CTGCGCTGCGACCGCTCCGTTTC.....AGCAAAATT 417
136 alGlyleuGlylAlaThrPheProThrHisValLysleuSerProPro 152
418 CCGCGCGTCGATGCGGAGCGCTTGGCCATCTTCGTCATCGGATGACAC 467
153 ProGlyAsnLysAlaGlu.....lleuLeileAsnAlaValGluC 167
468 CAATCCGCTGGCTGCCGACCTACGCTATTCATCAAGAAGCCGCGAGG 517
167 sgluProTYrleuThrSerAspHisValleuMetleuGluHisGlyIug 184
518 ATTTCAAACGCGCGCTGTGTGATTGAGCCGTTGACCGAAGCGCAAAATC 567
184 IuileMetlleGlyValSerlleuMet.....LysAlaIle 196
568 CATGTTGTAGGAGCT...GGCGGAGCGCTCGCTGAAAATGCT.. 612
197 GluValAsnLysAlaVallleGlyValGluAsnAsnLysAspAlaI 213
613 .....GCCAATCATCAACACATG 631
213 eAlaHisleuThrLysleuAlaThrAlaTYrProGlylleGluValMetP 230
632 AATTCGGCGCGCGCATCCCTGCGGTTGATGGCAGCAGCATTCATTTC 681
230 roleuLysValGluTYrProGluGlylGlyLysGluIleuIleAspAla 246
682 ATC.....GAGCGGCTGGCGCGGAATA 704
247 ValIleArgLysGluValLysSerGlyAlaLeuProIleSerThrGlyAl 263
705 AACCGGTGACCATCATTAAGATGTAATACCATGGCCGTTTGT 754
263 aValVal.....GlnAsnValGlyThrValPheAlaValI 275
755 TTGCAACAGCGCGTGAAC.....ACGAGCGCGGTGATGCCCTA 795
275 YrGluAlaValGluLysAsnLysProLeuValGluArglleValI 291
796 GGTGGTTCAGTCAACAACCGCGCTTGTGTCAGCTTGTGGTGC 845
292 ThrGlyLysleuSerArgProSerAsnLeuLeuValArgIleGlyTh 308
846 GAAAGATCGCAAAATT.....ACTGCGGCGCAATTGGTTGACACAGACA 889
308 rProIleAlaIleAlaLeuIleGluAlaIleGlyLysGlyLeuProGluAsnTh 325
890 ACCGCGGATTCGGGTGCGTATTCAGCGCGCGATTACACAGCGCG 939
325 LysIlelleGlyGlyLys..... 331
940 CACGATTATTGGAGCGCTACACATCAGATTTCGTTATGAGAAGAG 989
331 ..... 331
990 CCGCAGAAAGAGCTGTGCGGTGGGTGGCGGACCGGACAAATACT 1039
332 .....PrometMetGlyArgAlaLeuLeuSerProAsp...ValP 344

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1040 CCAFCACGCGTACACACCTCGCCATTTCCTGAAAAACAACACTTTCAAG 1089
344 roValThrLysGlySerSerGlyValleuIleLeuAspTrg..... 357
1090 TTCACACAGCGCGTCAACAGCGCGCGGACCGGCCCATGCTGCCATTGTAC 1139
358 ..GluGluAlaValAlaArgLysPrometArgAspCysIleArgCysAla 373
1140 TTACAGACGCGGTGATGCGCTTGGATATCCGCCACACCTGTTTGGCGG 1189
373 scYsValGlyValCysPrometGlyLeuAsnProAlaPheleuMetArgA 390
1190 ATTATATCGTGGCGCATACCGACAGCGCGCAG.....GCATTGGGT 1230
390 spThrleuTYrLysSerTrpGluThrAlaGluLysGlyAsnValAlaIsp 406
1231 TGCTTGGAATTGGAGACAGAGAACGCTGCTTGTGACGCTGTCGCCG 1280
407 CysIleGluCysGlySer.....CysSerPheThrCysP 418
1281 GCGCAAAATCGAATACGCGCGCTGTGGCGCAAGTGTGAAACATT 1329
418 oAlaAsnArgProLeuLeuAspTYrIleArgGlnAlaLysLysThrVal 434

seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA199.DAT:AA134343
seq_documentation_block:
ID AA134343 standard; Protein: 451 AA.
AA134343;
25-AUG-1999 (first entry)
Porphorymonas gingivalis protein Pg122.
XX DE Porphorymonas gingivalis; Pg; periodontal disease; gingivitis;
XX KW vaccine; antigenic.
XX OS Porphorymonas gingivalis.
XX PN W09929870-A1.
XX PD 17-JUN-1999.
XX PF 10-DEC-1998; 98WO-AU01023.
XX PR 04-AUG-1998; 98AU-0005028.
XX PR 10-DEC-1997; 97AU-0000839.
XX PR 31-DEC-1997; 97AU-0001182.
XX PR 30-JAN-1998; 98AU-0001546.
XX PR 10-MAR-1998; 98AU-0002264.
XX PR 09-APR-1998; 98AU-0002911.
XX PR 23-APR-1998; 98AU-0003128.
XX PR 05-MAY-1998; 98AU-0003338.
XX PR 22-MAY-1998; 98AU-0003654.
XX PR 29-JUL-1998; 98AU-0004917.
XX PA (CSLC-) CSL LTD.
XX PI Agius CT, Barr IG, Hocking DM, Margetts MB, Patterson MA;
XX PI Ross BC, Rothel LJ, Webb EA;
XX PT WPI; 1999-385613/32.
XX DR N-PSDB; AA191561.
XX PT Antigenic Porphorymonas gingivalis peptides for preventing
XX PT gingivitis
XX PS Claim 1; Page 303; 588pp; English.
XX PS AA191536 to AA191801 encode two hundred and sixty six antigenic
XX CC Porphorymonas gingivalis (Pg) polypeptide sequences given in AA191318 to

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CC AAY34583. AAY91802 to AAY91989 represent PCR primers used in the isolation of the PG polypeptides. The PG polypeptides have antibacterial activity with a vaccine mechanism of action. The PG polypeptides can be used as vaccines especially against Porphyromonas gingivalis. Probes can be used to detect Porphyromonas gingivalis in standard hybridisation assays. Porphyromonas gingivalis is involved in periodontal disease especially gingivitis.

XX Sequence 451 AA:

alignment_scores: Quality: 169.50 Length: 483
Ratio: 0.712 Gaps: 21
Percent Similarity: 49.275 Percent Identity: 20.290

alignment_block:
US-09-303-518D-125 x AAY34343 ..

Align seg 1/1 to: AAY34343 from: 1 to: 451

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37 GCGGCGACAGCCGAGCAAGCCGTTACAGCGCCGCCATTACCGAAGT 86
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27 AlaglyspProvalGluValLeu.....ProleproSerGlnva 40
87 CGCGTGTCTGGCGAAGATATGCCGGATGCCGCCCTCCGATGAAGTCA 136
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40 lValilleProleuglGlnHisileglyAlaProAlathrVal 57
137 AGGAAGGCGATGCGTCAAAAAGGCGCAAGTGTGTTGAGACAAAAG 186
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57 yslvsglyAspGluVallyValglyThrIlelle..... 68
187 AATCCGGCGGTGCTTACTGCGCGCGCTTACGCAAAATCGCCGGC 234
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69 .....AlaglnAlaglyGlyPheValSerAlaAs 78
235 .ATTACCGCTGGC...GAAAAGCGGCTCTCAG..... 264
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78 nIleHisSerSerValSerGlyValLeuLysIleAspAsnValIytrA 95
265 .....TCAGTGTGATTGCGGTGGAAGGCAAC 291
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
95 spserSerGlyTyrProLysProAlaValPheIleSerValGluGlyasp 111
292 GAGCAATCGAGTTGAAGCTACGCACTGAGCGCTG.....GCAAA 335
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
112 GlutPrGluGluGlyIleAspArgSerProAlaIleVallyGluCysAs 128
336 CTTAAGCGGCGAAGAGTGGCGCGCAACCTGATCCATCCGATTTGTGA 385
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
128 neuAspAlaLysGluIleValAlaLysIleSerAlaIleGlylle...V 144
386 CTGCGCTGGCGACCCGCTCCGCTTC.....AGCAAAATT 417
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144 alGlyLeuGlyGlyAlaThrPheProHisVallyLeuSerProPro 160
418 CTGCGCTGGAGTGGCGAGCCGCTGCGCATCTTCATGGATGGAAC 467
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161 ProGlyAsnLysIleGlu.....IleLeuIleIleAsnAlaValGlyC 175
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518 ATTTCAAGCGCGCTGTGTGATTGAGCGCTTGAACGCAAAATC 567
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568 CATGTTTGAAGCAAGCT...GGCGAGACGTCGCGTGAAGAAATGCT 612
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632 AATTCGGCGCGCGCATCTCGCGGTTGAGTGGCAGCAGCATTCATTC 681
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238 roLeuLysValGlnIlyrProGlnGlyGlyGluLysGlnLeuLeuAspAla 254
682 ATC.....GACGCGTGGCGCGCAATTA 704
255 ValIleArgLysGlnValLysSerGlyAlaLeuProIleSerThrGlyAl 271
705 AACCGTGTGACCATCATTAATCAAGATGTAATTAATTCATTCGCGCTGT 754
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300 ThrGlyLysLysLeuSerArgProSerAsnLeuValArgIleGlyTh 316
846 GAAAGTATCGCAATAT.....ACTGCGGCGCAATTTGTTGACAGACAGA 889
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316 rProIleAlaIleAlaLeuIleGluAlaIleGlyLeuProGluAsnThrG 333
890 ACCGCGTATTTCCGTTGCGTATTTGACGCGCGCATTCACAAAGCGCG 939
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333 LysIleIleGlyGlyGly..... 339
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339 ..... 339
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366 ...GluIleAlaValaLargLysPrometArgAspCysIleArgCysAlaL 381
1140 TTACGAGCGCGTATGCGCTTGGATATCTGCGCACCTGCTTTTCCGGC 1189
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381 scYsValGlyValCysPrometGlyLeuAsnProAlaPheLeuMetArg 398
1190 ATTTAATCGTCGCGCATACGACAGCGCGCAG.....GCATGGGT 1230
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415 CysIleGluLysGlySer.....CysSerPheThrCysPr 426
1281 GGGCAATATACGATACGCGCGCGCTGTGCGCAAGTCTGGAACATTC 1329
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426 oAlaAsnArgProLeuLeuAspTyrIleArgGlnAlaLysLysThrVal 1442

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seq_documentation_block:

ID AAB59813 standard; Protein; 1017 AA.

XX AAB59813;

XX 04-APR-2001 (first entry)

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436 rgleuSerAlaSerSerIleasMetArgSerThrSerValSerAlaPro 452
1124 ...TGTCGGCATTGCTACTACAGCGCGTGAATGCCCTTGATATCCTG 1170
453 ArgThrCysArgAlaThrSerSerSerAlaSerCysArgCysLeuSerCys 469
1171 CCCA.....CCCCGCTTTT 1184
469 sProGlnSerThrThrAlaAlaTrpAsnSerGlyTTrpThrProAlaProC 486
1185 GCGCGATTATATCGTCGCGCATACGACAGCGCGCGCAT.....1225
486 ySProSerSerProMetAlaGlyThrThrArgSerArgArgSerSerArg 502
1226TGGTTGCTTGGAATTTGACGACGAGAACGCTCGCTTG 1263
503 ArgThrProSerSerTrpProSerArgAsnTrpTrpSerArgArgAsnTh 519
1264 TGCAGCTTCGTCGTGCGCGCAATACGAAATACGACGCGCTGCGCAA 1313
519 rProSerSerAsnSerAlaLysArgArgGlyThrGlyLysValSerArgL 536
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536 yScys 537
seq.name: /SIDSI/gcgdata/geneseq/geneseq-emb1/AA2001.DAT: AAB59826.
seq_documentation_block:
ID AAB59826 standard; Protein; 1615 AA.
AC AAB59826;
XX
XX
XX AAB59826 (first entry)
DT 04-APR-2001
XX
XX Protein #3 encoded by TufD/E gene.
DE
XX
XX Toluene degradation; enzyme; waste degradation; TufD; TufD.
KM
XX
XX Thauera aromatica.
OS
XX Xanthomonas maltophilia.
OS
XX Geobacter metallireducens.
OS
XX Azococcus toluyticus.
XX
XX W0200072650-A2.
XX
XX 07-DEC-2000.
PD
XX
XX 24-MAY-2000; 2000WO-US14298.
PF
XX
XX 01-JUN-1999; 99US-0323872.
PR
XX
XX (UYOH-) UNITV OHIO.
PA
XX
XX Coschigano FW;
PI
XX
XX WPI: 2001-041080/05.
DR
XX N-PSDB; AAF23627.
DR
XX
XX
XX Composition comprising toluene degrading enzyme useful for biological
PT treatment of organic compounds, especially for degrading toluene or its
PT analogs
XX
XX
XX Disclosure; Fig 12; 122pp; English.
XX
XX The present invention relates to toluene degrading enzyme genes and
CC proteins tuch (see AAF23629 and AAB59831), tufI (AAF23630 and AAB59832),
CC tufF (AAF23631 and AAB59833) and tufG (AAF23632 and AAB59834). The
CC tufF (AAF23631 and AAB59833) and tufG (AAF23632 and AAB59834). The
CC toluene degrading enzymes are homologues of pyruvate formate lyase. The
CC toluene degrading enzymes are useful for biological treatment of organic
CC compounds and in particular for the degradation of toluene and its
CC analogs contained in liquid or solid waste source. The present sequence
CC is a protein sequence encoded by toluene degrading enzyme gene, TufD/E.

XX
SQ Sequence 1615 AA:
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Quality: 166.50 Length: 486
Ratio: 0.733 Gaps: 26
Percent Similarity: 46.708 Percent Identity: 25.103
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103 GAATATGCGGTATGCGCCCTCGATGAAAGTCAAGAGAGCGATCGGT 152
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723 gArgValSerThrThrSerProArg..SerThrGlyArgArgTrpSer 739
153 CAAAAAGGCCAAGTCTGTTGAAGACAAAAAGATCGGCGGTGT 202
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740 SerProAlaArgArgSerAlaGlyArgAlaGlyArgAlaGly..... 753
203 TTACTGCGCGGCTTCAGGCAAAATGCGCGATTC.....CGTGGCGAA 249
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754 ...CysAla.....ArgSerArgGlyThrSerArgProIleArgS 767
250 AAGCGCGT.....ACTCAGTCAGT..... 269
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767 eraLarArgProSerCysSerLysSerProThrSerValSerAlaPhePro 783
270CGTATGCGGTGCAAGCAACGACGAAATCGAGTTG 307
784 ProSerProAlaArgAlaSerArgThrArgCysArgArgAsnSerLeuPr 800
308 AAGCTACGACCTGAGCGCTGCGCAAACTTAAGCGCGCAAGATGCGC 357
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800 oSerSerValThrArgSerSerAla.....ThrArgAlaAlaThrP 814
358 CGCAACCTGATCCAAATCGGTTGTGACTGCGCTGCGACCCGTCGTT 407
|||: |||: |||: |||: |||: |||:
814 roArgArg..LysThrProCysCysGlyArgThrThrArgProPheSerS 830
408 CAGCAAAATTCCTGCGGTGATGCGCGCTTCGCACTTCGTCGAAG 457
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830 erThrArg.....AsnSerSerArgAlaThrTrpMet 840
458 CGATGACACCAATCCGCTG.....CTGCGACCCCTACGGTCATT 498
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841 ArgTrpAsnSerSerArgTrpAsnValArgPheProSerMetAlaProAl 857
499 ATCAAGAGAGCGCGCGAGATTTCAAAGCGCGCTGTGATATGACCG 548
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549 TTGACCGACGCAAAATCATGTTGTAAAGCAGCTGGCGGCGACGTCG 598
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871 leCysSerSerSerProSerAlaAlaProThrProArgAlaArgThrPro 887
599 CGTCTGAAATGCTGCCAATCAACATGAAATTCGCGCGCGCAT 648
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888 AlaThrThrProThrProSerSerArgGlnProSerGlySerLarArgPr 904
649 CTGCGCGTTGATGTCGACGACGACATTCATTTCATGACG..... 688
904 oSerProPro.....SerSerSerAlaIleProA 914
689 ..CGTGGCGCGCAATTAACGTTGGACCAATTCATTCAGATGTAA 736
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914 rGArgThrAlaArgArgArgCysAlaGly..... 923


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110 ArgGlyProThrAlaAlaArgSerGlyArgHisGlyArgArgGlyTrpArg 126
282 .....TGAAAGCAGACGACCAATCGAGTTTGAAACGCTACGACACT 321
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322 GAAGGGCTGGCAAACTTAAGGGGGGAGAGAGTGGCGGCAACCTGATCA 371
136 .....ArgArgArg..... 138
372 ATCCGGTTTGTGACTGCGCTGCCACCCGCTCCGTTGACGAAATTCCTG 421
138 ..... 138
422 CCGTCGATGCCGAGCCGTCGCCATCTTCGCAATGCGATGACACCAAT 471
139 .ArgArgCysArgArgValArgArg.....GlyArgArg 150
472 CCGCTGGCTGGCCGCTACGCTGATTAAGAGAGAGCCGCGAGATTT 521
150 eArgArgCysArgArgGlnArgHisAlaAspArgAlaArgArgArg 166
522 CAAGCGCGCTGTGTATTGAGCCGTTTGACGCAAGCAAAATCCATG 571
167 ArgArgArg.....ArgArgGlnGln..... 173
572 TTTGTAAGGACGCTGGCCAGACGTCGCTGAATAATGC.....TGC 615
174 ...PheArg**TrpArgGlyArgArgArgGlyArgCysArgArgAla 189
616 AACATCGAAGACATGAATTCGCGCGCCGATCCTGCGGTTTGAGTGG 665
189 rOlenuArgGlnTrpArgValArgArgProArgArgSerArgHis..... 203
666 CAGCAGCATTCATTCATCGAGCCGCTGGCGCGCAATAAACCCTGTGGA 715
204 .....GlyArgGlnHis..... 207
716 CCATCAATTATCAATGATTAATACCATTTGGCCGTTTGTTCACACAGC 765
208 .....ArgTrpArgTrpArg.....ArgArg 215
766 CGTCTGAACACCGACGCGGTATGCGCTAGGTGTTCTCAAGTCACAA 815
215 rArgTrpArgGlnAlaAspArgProArgTrp..... 226
816 ACCGGCCCTTTCGCTACCGCTTTGGGTGCGAAGTATCGCAATTACTG 865
227 .....ArgArgArgCys..... 230
866 CCGGGGAATTTGGTGTACACAGCAACCGGTGATTTCCGGTTCGGTATTG 915
231 .....ArgArgGlyProArgArgSerLeuGlyTrpPro* 242
916 AACGGCGCATTTACACAGGCGCGACGATTTATTTGGAGCTACACAGA 965
242 **ArgArgAlaArgTrpArgGlnArgTrpArgProAlaIleArgGln 258
966 TCAGATTTCCGTTATCGAAGAGCGCGACGCAAGA..... 1001
259 .....ArgArgArgArgArgProArgHisArgArgAsnThrAlaG 272
1002 ....GCTTGGCGGTGGGTG.....GCCGAGCGC 1028
272 yGlyGlyGlnArgGlyIleGlyAspGlyPheValArgCysTrpArgPro 289
1029 GGAACAAATATCTCATCGACGCTACAAACCTCGGCATTTCTGAAGAA 1078
289 rArgHisArgProThrArgLeuAlaPro.....Ile 299
1079 AACTTTCAAGTTCAACACAGCCG.....TCACAGGC 1110
300 AsnGlnGlyPheGlyAlaGlyProGlnHisGlyHisProLeuSerTrpArg 316

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327 .....ProGlnGlyProLeuArgLeuGlyGlnPheLeuAla 339
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1261 TTGTGCGAGTTGCTGTCGCCGCAATACGATACG...GCCGCTGT 1307
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361 ys**AlaCysTrpLeuPro 367

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ID AAB59827 standard; Protein: 1592 AA.
AC AAB59827;
XX
XX
XX 04-APR-2001 (first entry)
XX
XX Protein #4 encoded by TufD/E gene.
XX
XX Toluene degradation; enzyme: waste degradation; TufE; TufD.
XX
XX Thauera aromatica.
XX
XX Xanthomonas maltophilia.
XX
XX Geobacter metallireducens.
XX
XX Azococcus toluylicus.
XX
XX W0200072650-A2.
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XX 07-DEC-2000.
XX
XX 24-MAY-2000; 2000WO-US14298.
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XX 01-JUN-1999; 99US-0323872.
XX
XX (UYOH-) UNIV OHIO.
XX
XX PA
XX
XX Coschigano PW;
XX
XX WPT: 2001-041080/05.
XX
XX N-PSDB: AAF23627.
XX
XX Composition comprising toluene degrading enzyme useful for biological
XX treatment of organic compounds, especially for degrading toluene or its
XX analogs
XX
XX Disclosure; Fig 12; 122pp; English.
XX
XX The present invention relates to toluene degrading enzyme genes and
XX CC proteins tufH (see AAF23629 and AAB59831), tufI (AAF23630 and AAB59832),
XX CC tufF (AAF23631 and AAB59833) and tufG (AAF23632 and AAB59834). The
XX CC toluene degrading enzymes are homologues of pyruvate formate lyase. The
XX CC toluene degrading enzymes are useful for biological treatment of organic
XX CC compounds and in particular for the degradation of toluene and its
XX CC analogs contained in liquid or solid waste source. The present sequence
XX CC is a protein sequence encoded by toluene degrading enzyme gene, TufD/E.
XX
XX Sequence 1592 AA;

```

alignment_scores:

Quality: 146.00 Length: 527
 Ratio: 0.705 Gaps: 31
 Percent Similarity: 39.279 Percent Identity: 23.909

alignment_block:

US-09-303-518d-125 x AAB59827

Align seg 1/1 to: AAB59827 from: 1 to: 1592

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18 AGSTCTAAACCTGCCCATCGCGGAGACCGGACACCGTTTA..... 62
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
689 ArgAlaIaIaProIaIaCysAlaIaArgPheAlaIaTyrArgArgLeuGlyPr 705
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
63 .CGAGCGCCCGGCGCATTCACGAGTGGCGTTCGGGAGAAATATATCC 111
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
705 OArgGlyThrSerArgGlyArgSerArgCysSerProAspArgCysCysA 722
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
722 IgrTrpSerArgCysSerSerProSerProArgArgCysProProSerSer 738
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
162 CCAAGTGGCTGTTGAAGCAAAAGCAATCCGGCGGTGTGTACTGCGC 211
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
739 ProIaIa.....GlyIaIaProGlyIaIaThrCys.. 747
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
212 CGGCTTCAGGCAAAATCCCGCGCATTCACCGTGGCAAAACGGGTACTT 261
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
748 .....SerArgArg.....ProPheSerArgSerArgAspS 758
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
262 CAGTGAAGT..CGTATGCGCGTGAAGGACAGCAAGCAATGAGTTTGA 308
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
758 erAlaGlyProIaIaArgPheArgArgCysArgAsp..... 771
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
309 ACGTACGACACCTGAAGCGCTGGCAACTTAAGCGGAGAAAGATGGGCG 358
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
772 .....AlaCysGlyIaIaArgAlaIaArgCysPr 780
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
359 GCAACTGATCCAAATCCGTTTGTGAGTCGCGTCCGACCCGTCGCTTC 408
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
780 OGlyPro.....ArgSerAlaProSerIleA 789
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
409 AGCAAAATCTCGCGCGATGCGGACCGCTGCGC..... 443
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
789 rgrArgGlySerArgAspArgSerArgAlaSerArgSerArgGlySerPro 805
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
444 .....CATCTGTCATATGCGATG 463
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
806 LeuGlyAlaIaIaThrSerCysProArgArgArgArgCysSerIle 822
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
464 ACACCAATCCGCTGCGCGACCGCTACGGTCAATTATCAAGAAGC.... 509
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
822 eGlyIaIaSerSerGlyCys...ProHisProIaIaArgArgSerProV 838
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
510 .....CGCCGAGGATTT 521
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
838 alaIaIaSerSerLeuArgAlaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 853
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522 CAAACGCGCGCTGTGGTATGAGCCGTTGACCGAGCAAGCAAAATCATG 571
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
854 .....ArgPheArgGlyProIaIaIaIaIaIaIaIaIaIaIaIaIaIa 868
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
572 TTTTGAAGGACGTCGGCGGACAGTGGCTGTGAAATGC...TGCCAC 618
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
868 s.....TrrArgTrpProArgProArgArgCysArgCysSerA 881
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619 ATCGAAACACATGAAATCGCGCGGCC.....GCATCTGCG...CGG 656
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
881 rgrArg.....TrpGlyArgProIaIaIaIaIaIaIaIaIaIaIaIaIaIa 894
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
657 TTTGAGTGGAC.....GC 670
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
895 AlaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 911
   ||||| ||||| ||||| ||||| ||||| ||||| |||||

```

```

671 ACATTCAATTTCATGAGCCGCTGCGGCCGCAATTAACCGTGGACCATC 720
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
911 aserProIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 928
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
721 AATTATCAAGATGTAAATTACCATTTGGCCGCTTTGTTGCAACAGCCGCT 770
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
928 rgrArgSerArgCys.....ProIaIaSerSerProIle 938
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
771 GAACACCGGAGCGGTATGTCCTAGTGGTTCCTCAATCAACAAACCCG 820
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
939 ArgTrpThrGlyIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 949
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
821 GCCTTTGCGCTACCGTTTGGCTGGCAAGTATCGCAAT.....T 861
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
949 gProIaIa.....GlyCysSerProIaIaIaIaIaIaIaIaIaIaIaIaIa 962
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
862 ACTCGCGGCAATTTGCT.....TGACACAGA 887
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962 rGlyCysGlyIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 978
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
888 CAACCGCGT.....GATTTCCGGTTCGTTATTTGAACGCGC 922
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
979 SerIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 994
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
923 CGATTACAGAGGCGGCGAGCAATTTATTTGGAGCGGACCAATCAGATT 972
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
994 gSerThrThrIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1011
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
973 TCCGTTATCGAAGGCGGCGAGCAAAAGAGCTGTTCGGCTGGGTTGCGCC 1022
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1011 hIaIaSerThrIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1025
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1023 .....GCAGCCGAC 1033
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1026 ThrAspIleHisSerGlyIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1042
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1034 AATATCTCATACAGCGGTACACCTCGGCCATTTCTGTAACAAACATC 1083
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1042 gAlaIaIaSerGlyIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1059
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1084 TTCAGTTCACACAGCGGTCAACG..... 1109
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1059 ysProIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1075
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1110 .....CGGCGACCGCGCCATGGTGGCGATTGTTACTT 1141
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1076 AlaCysGlySerSerSerArgArgProSerSerGlyIaIaIaIaIaIaIaIa 1092
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1142 ACGAGCGGTGAT.....GCCCTGGATAT..... 1166
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1092 IProIleArgProSerSerIleCysGlyArgAlaIaIaIaIaIaIaIaIaIa 1109
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1167 .....CTGGCCAC 1175
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1109 roSerSerProIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1125
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1176 CCGCTTTTGGCGGATTAATCGTCGGCATACCGACAGCGCGAGGAT 1225
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1126 ThrProCysArgArgHisIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1136
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1226 TGGGTTGCTTGGAAATTTGACGAGAAGACCT 1256
   ||||| ||||| ||||| ||||| ||||| ||||| |||||
1137 .....GlySerArgArgPro 1141
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seq_name: /SISL/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAB59817

seq_documentation_block:

ID AAB59817 standard; Protein: 999 AA.

XX AAB59817;

XX

DT 04-APR-2001 (first entry)

XX

DE Tutd protein #8.
 KM Toluene degradation; enzyme; waste degradation; Tutd.
 XX
 XX Thauera aromatica.
 OS Xanthomonas maltophilia.
 OS Geobacter metallireducens.
 OS Acetobacter toluolyticus.
 XX
 XX WO200072650-A2.
 PN
 XX 07-DEC-2000.
 PD
 XX 24-MAY-2000; 2000MO-US14298.
 PF
 XX 01-JUN-1999; 9905-0323872.
 PR
 XX (UYOH-) UNIV OHIO.
 PA
 XX Coschignano PW;
 PI
 XX WPI, 2001-041080/05.
 DR N-PSDB; AAF23625, AAF23627.
 XX
 XX Composition comprising toluene degrading enzyme useful for biological
 PT treatment of organic compounds, especially for degrading toluene or its
 PT analogs
 XX
 XX Disclosure; Fig 5; 122pp; English.
 PS
 XX The present invention relates to toluene degrading enzyme genes and
 CC proteins tutd (see AAF23629 and AAB59831), tuti (AAF23630 and AAB59832),
 CC tutf (AAF23631 and AAB59833) and tutg (AAF23632 and AAB59834). The
 CC toluene degrading enzymes are homologues of pyruvate formate lyase. The
 CC toluene degrading enzymes are useful for biological treatment of organic
 CC compounds and in particular for the degradation of toluene and its
 CC analogs contained in liquid or solid waste source. The present sequence
 CC is a protein sequence for toluene degrading enzyme, Tutd.
 XX
 SQ Sequence 999 AA;

alignment_scores:
 Quality: 145.00 Length: 529
 Ratio: 0.700 Gaps: 31
 Percent Similarity: 39.130 Percent Identity: 23.819

alignment_block:
 US-09-303-518d-125 x AAB59817 ..

Align seg 1/1 to: AAB59817 from: 1 to: 999

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18 AGGTCTAAACCTGCGCCATCGCGGAGACCGAGCAAGCCGTTTAA..... 62
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94 ArgAlaIaIaProAlaCysAlaArgArgPheAlaTyrArgArgLeuGlyPr 110
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
63 CGAGCGCGCGCGCATACGGAAGTCGGTGGTGGCGGAGAGAAATATGCC 111
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
110 ArgGlyCysThrSerArgGlyArgSerArgCysSerProAspArgCysCys 127
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
112 GGATATCGCGCCCTCGATGAAGTCAAGAGAGCGATGCCGTCAAAAAAG 161
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
127 rgrPrSerArgCysSerSerProAspArgArgCysProProSerSer 143
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
162 CCAAGTGTGTTTGAAGACAAAAAGATCCGGCGGTGCTTACTGCGC 211
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
144 ProAla.....GlyAlaProGlyAlaThrCys.. 152
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212 CGGCTTCAGGCAAAATCGCGGATTCACCGTGGCGAAAGCGGACTT 261
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153 .....SerArgArg.....ProPheSerArgSerArgAsp 163
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262 CAGTCACT..CGTGAATGCCGTTGAAGCAACGAGCAAAATCGAGTTGA 308

```

```

163 erAlaGlyProAlaGlyAlaAlaArgPheArgArgCysArgAsp..... 176
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309 AGCCTACGCAACCTGAAGCGCTGGCAAACTTAAGCGCGGAGAGAGCGGC 358
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
177 .....AlaCysGluAlaGlyAlaArgCysPr 185
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
359 GCACCTGATTCNAATCGGTTGTGTGAGTCGCGTCCGACCGCGTCC 408
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185 OGlyPro.....ArgSerAlaProSerIleA 194
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
409 AGCAAAATTCCTGCGCTCGATGCGGAGCGGTTGCG..... 443
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
194 rgrArgGlySerArgAspArgSerArgAlaSerArgSerArgArgGly 210
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
444 .....CATCTGCTCAATG 457
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
211 SerProLeuGlyGlyAlaThrAlaThrSerCysProArgArgArgArgCys 227
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
458 CGATGACACCAATCCGCTGGCTGGCGGACCGCTACGCTCATTTCAAGAA 507
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
227 sSerIleGlyAlaSerSerIleCys..ProHisProValArgArgS 243
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
508 GC.....CGCCGA 515
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
243 erProValaIaSerSerIleArgAlaHisArgGlyCysThrAlaArgArg 259
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
516 GSATTTCAAGCGCGGCTGTGGTATTCAGCCGTTGACGAGCAAGCAAA 565
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
260 Gly.....ArgPheArgGlyProThrSerArgAspThrIleArgArg 273
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
566 TCCATGTTTGTAAAGCAGCTGGCGGACAGCTGCCGCTGCAAAATGC...T 612
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
273 gArgCys.....TrpArgTrpProArgProArgArgCysArgC 286
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
613 GCCAACATTCGAACACATGATTCGGCGGCC.....GATCCTCG 653
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
286 ySerArgArg.....TrpGlyArgProLeuThrAlaSerGlyCys 299
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
654 ..CGTTTGAAGTGGCAC..... 668
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
300 ProArgAlaArgTrpArgArgGlySerAsnTrpSerSerIleArgSerSe 316
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
669 .....GCACATTCATTTTCATCGAGCGCGGTCGCGGAAATTAACCGTGTG 714
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316 rAlaAlaSerProLysArgThrCysGlyArgGlyValArgSerAspHis 333
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
715 ACATTCATTAATTCAGATGTAATTAATTCATTCGCGCTTTGTTGCACAGG 764
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
333 erAlaArgArgSerArgCys.....ProAlaSerSer 343
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765 CCGTCTGAACACGAGCGCGGATTCGCTAGGTGCTTCAGATGTAACA 814
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
344 ProLeuArgTrpThrGlyArg...CysArgArgTrp..... 354
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815 AACCGCGCTCTGCTGATACGCTTTGGTGGCGAAGATATGCAAT... 860
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
355 ArgArgProLeu.....GlyCysSerProArgAlaThrCysT 367
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
861 .....TACTGGCGGCGAATGTG.....TGA 881
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367 hrAlaArgCysGlyArgAspGlyCysSerAlaPhePheGlyAsnProLeu 383
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
882 CACAGACAAACCGCT.....GATTCGCTGGTGGTATGA 916
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384 HisArgSerLeuArgGlyProThrAlaAlaProPheArg...AlaHisAr 399
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
917 ACGGCGGATTAACAAAGCGCGACGATTAATTTGGACCGTACCAAT 966
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
399 gSerArgSerThrThrArgArgCysAlaValArgIleSerSerArgHisA 416
   |||:|:| |||:|:| |||:|:| |||:|:| |||:|:|
967 CAGATTCGCTTATCGAAGAGCGCGCAAAAGCTGTCGGCTGGGT 1016
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416 spArgThrAlaSerThrArgArgProHisLysProPro.....LysGly 430
1017 TGCGCC.....GCGGC 1027
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431 CysAlaIleThrAspIleHisSerGlyArgGlyCysTrpProAlaThrAlaSe 447
1028 CGGAAATAACTCCATCAACCGGTACACCGGTGGCCATTCCGTGAAAG 1077.
|||||
447 IserArgAlaAlaSerGlyAlaSerAlaLysArgThrArgLeuArgArgA 464
1078 AAACCTTCAGTTCACACAGCGGTCAACG..... 1109
|||||
464 rGserCysProValArgSerProAlaArgArgGlyThrArgAlaAlaTrp 480
1110 .....CGCGACCGCGCCATGGTGGCGGATG 1135
|||||
481 HisSerAlaCysGlySerSerSerArgArgProSerSerGlyAlaArgProTr 497
1136 GTACTTACGACGCGGTGAT.....GCCCTTGAGAT. 1166
|||||
497 pSerValProIleArgProSerSerIleCysGlyArgAlaValGlyLeuT 514
1167 .....CCT 1169
514 hrserProSerSerProLeuAsnArgProPheAlaArgArgSerAlaPro 530
1170 GCCCACCCGCTGTTTGCAGATTAAATCGTCGCGGATACCGACAGCGCGC 1219
|||||
531 AlaSerThrProCysArgArgHisAsnArgArgArgTyr..... 543
1220 AGGCATTGGTGGCTTGGAAATTGACAGCAAGAACCT 1236
544 .....GlySerArgArgPro 548

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seq_documentation block:
ID ABG03569 standard; Protein; 819 AA.
XX
AC ABG03569;
XX
DT 13-FEB-2002 (first entry)
XX
DE Novel human diagnostic protein #3560.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
PI
DR WPI; 2001-639362/73.
DR N-PSDB; AAS67756.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity
XX
PS Claim 20; SEQ ID NO 33928; 103pp; English.
XX

```

CC The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. ABG00010-ABG30377 represent novel human
CC diagnostic amino acid sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

SO Sequence 819 AA;

alignment_scores:
 Quality: 125.50 Length: 388
 Ratio: 0.860 Gaps: 26
 Percent Similarity: 37.629 Percent Identity: 25.515

alignment_block:
 US-09-303-518D-125 x ABG03569 ..

Align seg 1/1 to: ABG03569 from: 1 to: 819

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328 CysProAlaLysArgGlyGlnProGlyCysGly***AlaProTr 342
403 CCGTTCAGCAAAATTCCTGCGCGTGCATGCCAGCGGTCCGCACTTGGT 452
|||||
342 pArgPro.....LeuProArgArgProSerSerValProProProA 356
453 CAATGGGATGACACCAATCCGCTGCTGCGGACCCCTACGGTCATTATCA 502
::|
356 la.....TrpSerProProGlnAspLeuProLeuGlySerLeuPro 370
503 AAGAAGCCGCCGAGATTTCAAACCGCGCTGTGGTATGAGCCGTT. 551
|||||
371 AlaLysProThrAsnGlyGlyProAlaLeuCysPhe.ProProProHis 387
552 .....GACGAGACCAAAAT..... 566
387 erLeuGlnProGlnAspAlaSerGlyLysThrGlnGlyProGluGluA 403
567 .....CCATGTTTGTAAAGCAGCTG..... 587
404 ProProProCysLeuValProArgTrpProProAspSerAsnSerArg** 420
588 .....CGCAGACGTGCGCTGTGAAATGCT..... 612
420 *HisProArgArgSer.ProMetSerProAlaProHisSerThrProGly 436
613 .....GCCAACAATGGAACACATGAAATC..... 636
437 ArgArgHisLeuThrGlnIleProHisnTrpLysThrHisLeuPhePro** 453
637 .....GGCGGCCGCACTCTGCC..... 654
453 *AlaProAlaArgGlyProSerProGlyArgAlaCysThrSerProCysp 470
655 .....GGTTTG.....AGTGGC 666
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1181 AGCAGGGTGGGAGATATCAAGAGGCATCAGCGCTGTAGTA...CC 1135
140 AsnProThrProValIleAlaIleGlySerThrProSerValAlaGlyPro 156
1134 AATGGCACCATTGGCGGGTCCCGCCGTTGAGCGCT... 1098
156 OleuGly...ValAsnSerProLeuSerAlaLeuGlyPheL 171
1098 ..... 1098
171 euThrSerAsnSphrAsnLeuIleAsnSerSerAlaLeuSerSerAla 187
1097 .....GTGTTGAACCTTGAAGATTGTTTTCAGGAATGCGC 1060
188 ValThrSerGlyLeuAlaSerLeuSerSerLeuThrLeuGlnAsn... 202
1059 GAGGGTGTACCGCTGATGAGATATTGTCGGCGTCCGCGCCACCCAGC 1010
203 .....SerAspSerSerAlaSerAlaP 210
1009 CGAACAGCTCTTTCCTGCGCGCTTCTCGATACGAATATCGATTGTGG 960
210 roAsn...LysCysTyrAlaProSerAlaIlePro..... 220
959 TAGCCTCCCAATTAATCGTGGCGCTTGTATCGCGCGCTTCATATC 910
221 ...ThrProGlnArgThrSerThrPro...GlyLeuAlaLeuPheProG 235
909 CGAACCGGAA.....ATCAGCGGCTTCTGTGTGCA 878
235 yProProSerProValAlaAsnSerThrSerThrProLeuThrLeuPro 252
877 CCAATTGCGCC.....GCAGTAATTTGCGATCTTTCGACCCAAA 837
252 aGlnSerProLeuAlaThrAlaAlaSerAlaSerThrSerAlaProVal 268
836 ACGGTACCCAGAGAGCGGCTTGTGACTTGAAGAACCCACTAGGCAAT 787
269 SerCysGlySerSerAlaSerLeuLeuArgLysProHisProGlyThr 285
786 CACGGCGCTGCTGTCAGACGCGCTTTCGAACAACAGCGCATGTAA 737
285 rAspLeuHisIleSerSerThrProAlaAlaThrThrLeuPro..... 299
736 TTACATCTTGAATTAATGATGCTCCACAGCGTTTATTCGCGCCAGCGC 687
300 .....ValMetIleLysThrGluProThrSerProThrPro 311
686 TCGATGAATGAATGTGCTGCCACATAA...CCGCGAGATGCGGGCC 640
312 Ser.....AlaPheLysGlyProSerHisSerGlyAs 322
639 GCCGAAT.....TCATGTGTTGCGATGTGCGAGCATTTTCAG 602
322 nProSerHisGlyThrLeuGlyLeuSerGlyThrLeuGlyAlaAlaTyrT 339
601 ACGGCAGCTGTCGCGCAGCTGCTTACAAACATGAGATTTCGCTCGTC 552
339 hrSerThrSerValProIleSerLeuSerAlaCysLeuAsnProAlaLeu 355
551 AAAGCGCTCAATACCAACAGCGCGCTTGA...TCTCGCGCGCTTC 505
356 SerGlyLeuSerSerSerSerThrProLeuAsnGlySerAsnProLeuSe 372
504 TTGATATGACCGTAGGTGCGAGCAGCGGATGTCATGCGCAT 455
372 rSerIleSerLeuProProHisGlySerSerThrProIleAlaProValP 389
454 TGACGAGATGGGAGACGGCTCGCATCGACGCGAGAAATTTGCTGAC 405
389 heThrAlaLeuProSerPheThrSerLeuThrAsnAsnProPheProLeuThr 405

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404 GAGCG...GTGCGACCGCAGTCCACAAACCGGATTGATC... 366
406 GlyAsnProSerLeuAsnProSerValSerLeuProGlySerLeuLea 422
365 .....AGGTTGCGGCGCACTTCTTCGCGCTTAAGTTTGCACGCGCT 323
422 aThrSerSerThrAlaAlaThrSerThrSerLeuProHisProSerSerT 439
439 hrAlaAlaValaLeuSerGlyLeuSerAlaSerAlaProValSerAlaAla 455
272 ACGACTGACTGAAGTAGCGCTTTCGCAAGTGAATGAGCGGCGATT 223
456 Pro.....PheProLeuAsnLeuSerThrAlaVala 465
222 GCCTGAAGCGCGCGCAGTAACACACAGCGCGGATCTTTCCTCA 173
465 lProSerLeuPheSerVal.....ThrGlnGlyProLeuSerSerSerA 480
172 ACAGCACTTGCGCTTTTTCAGCGCATCG...CCTTCCTTGACTTTC 129
480 snLeuSerTyrProGlyPheSerValSerAsnThrProSerValThrPro 496
128 ATCAGAGGGCGCATACCGCATATCTTCGCCAAGCAGCGCATTCGCT 79
497 AlaLeuProSerPheProGlyLeuGlnAlaProSerThrValAlaAlaVala 513
78 AATGCGCGGCGCG...TCGTAACGGCTTTCCTCGCTTCGCGCGGATG 32
513 lThrProLeuProValAlaAlaAlaThrAlaProSerProAlaProValLeuP 530
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530 roGlyPhe 532

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seq_documentation_block:
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XX
AC_ABB64198;
XX
DT_26-MAR-2002 (first entry)
XX
DE_Drosophila melanogaster polypeptide SEQ ID NO 19386.
XX
KW_Drosophila; developmental biology; cell signalling; insecticide;
XX pharmaceutical.
XX
OS_Drosophila melanogaster.
XX
PN_WO200171042-A2.
XX
27-SEP-2001.
XX
23-MAR-2001; 2001WO-US09231.
XX
PF_23-MAR-2000; 2000US-191637P.
XX
PR_11-JUL-2000; 2000US-0614150.
XX
PA_(PEKE ) PE CORP NY.
XX
PI_Venter JC, Adams M, Li PWD, Myers EW;
XX
WP1; 2001-656860/75.
XX
DR_N-PSDB; ABL0830L.
XX
PT genes from Drosophila and for elucidating cell signalling and cell-cell
XX interactions -
XX
PS Disclosure; SEQ ID NO 19386; 21pp + Sequence Listing; English.

```

XX The invention relates to an isolated nucleic acid detection reagent
 CC capable of detecting 1000 or more genes from *Drosophila*. The invention is
 CC useful in developmental biology and in elucidating cell signaling and
 CC cell-cell interactions in higher eukaryotes for the development of
 CC insecticides, therapeutics and pharmaceutical drugs. The invention
 CC discloses genomic DNA sequences (AB16176-AB16177), expressed DNA
 CC sequences (AB16177-AB16178) and the encoded proteins
 CC (AB16177-AB16178).
 CC The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 2406 AA:

alignment_scores:
 Quality: 119.50 Length: 492
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 Percent Similarity: 47.561 Percent Identity: 21.951

alignment_block:
 US-09-303-518D-125/rev x AB16178 ..

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 1294 HisArgSerSerAspSerArg..AsnSerArgGluSerProAlaSerL 1310
 1240 ATTCCAGCAACCCCAATGCTGCGCGCTGTCGATCG.....CGCAGC 1197
 :
 1310 euLysSerThrProSerAsnIleGlyLeuAsnValSerMetAlaProThr 1326
 1196 ATTAATG.....CGCAAAAGCAGGCGGCGACGATATCCAAAGCGAT 1153
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 1327 LeuArgSerIleThrProLeuAsnAsnSerSerAlaIleSerSerGlyAl 1343
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 1343 aserGln.....ProValSerValValProSerAlaAsnSerL 1357
 1102 CGCGTGTGTGACTTGAAGACTTTGTTTTCAGAGAAATGCCCGAGGTT 1053
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 1357 hVala...LeuSerMetSerAsn.....ProHisIle 1366
 1052 GThA.....CGCGTGAATGAGTATTTGTCGCGC..... 1026
 1367 SerHisSerHisValProAlaThrAlaSerGlyAlaPheSerSerSe 1383
 1025TCGCGCGCAACCCAGCCAGCACTTTG.....CTGCGGC 989
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 1383 rAlaAlaIaGlyThrSerThrProAsnSerGlyLeuSerThrLeuAlaV 1400
 988 CTTCCTTCGTAACGGAATCTGATGTGTACCGTCCCAATATATCTGCC 939
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 1400 alThrSerLeuSerThr.....SerAla 1407
 938 GCGCCTTGTAATGCGCGCTTCAATACGAGCAAGCAATCAGCGGTT 889
 ||||| :
 1408 AlaPro.....GlnProHisSerHisPheProGlnSerThrGlnMe 1421
 888 G.....TCGTGTG..... 882
 1421 tLeuProGlnSerGlyAsnPheSerSerValSerHisLeuThrThrN 1438
 881TCACCAATTCGCGCGAGTATTTGCGATACCTTTCCGA 843
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 1438 isPheMetSerGlnAsnGlnProMetValAlaGlyGlySer..... 1452

842 CCCAAACGGTACGACAGAGCGCGTTTGTGACTTGAAACCACTAG 793
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 1453ThrLeuLysSerIleSerSerAlaAlaIaThrAlaProProSe 1467
 792 GGCATACAGCGCGCTGCTGTCAGACGGCTTTCGCAACAAACGGCCAA 743
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 1467 rAlaAlaIa.....AlaValSerAsnPheThrProS 1478
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 1480 1480
 642 GCCCGCAATTCATGTTGATGTTGCGACCATTTTCAGACGCGCGT 593
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 1481AlaValGlnSerLeuThrAlaValThrSerSerSers 1494
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 1494 erSerProSerThrLeuSerSerSerValIleGlnLysValIleSerPro 1510
 551AAACGCTCAATACCAACAGCGCGCTTGAATTCCTGCGC 511
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 510 GCGCTCTTGATATAGCAGTACGAGTGGCGAGCGACGCGGATGTTGTCGA 461
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 1527 oAlaAsnAlaValValThrCysAlaProIleThrProIleValSer. 1543
 460 TCCGATTGACAGATGGCGAAGCGCTCGGACATCGACGCGAGGAATTG 411
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 1544SerGlySerAlaArgProThrProProLeu 1553
 410 CTGAAC.....GGACGGGTGCGGACGCGACGCTGCACAA 379
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 1554 SerAsnCysThrSerMetGlyIleGlyMetValAsnAlaIa..SerThra 1570
 378 ACCGATGATGATAGCTGCGCGGACCTTCTTGGCGCTTAAGTTGCCA 329
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 1570 laArg.....SerSerCysAsnAlaIle..SerProLeuSerIlePro 1584
 328 GCGCTTCAGT.....CGGTAGCGTTCAAC 303
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 1584 laThrAlaGlyIleHisValSerAlaThrAsnProSerPheGlnSerSer 1600
 302 TCGATTTGTCGTGCTGCT..... 285
 1601 SerTyrPheProThrProLeuAlaProProProSerSerProSerProAl 1617
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 1617 aThrSerSerIleAlaIleIleSerSerSerAlaSerGlnPheAsnPro 1634
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 200 ACCACGCGCGGATTTCTTTTGTCTTCAACACGACTTGGCTTTTGGAC 151
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 1651 ThrThr.....AlaSerSerValThr..... 1657
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 1673 isProPheSerAlaGluSerLeu 1680


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967 .AsnAsnAspLeuGlyAlaGlyMetAlaValAlaMetLysAspLeuGluM 983
809 TCACAAACCGCCGCTTGGCTACCGCTTTGGTGGCAAGTATGCCAA 858
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983 et.....ArgGlyAlaGlyAsnValLeuGlyAlaGluGlnSerGly 996
859 ATTACTGCG.....GGCGAATT 875
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997 HisIleAlaGlyValGlyPheAspLeuTyrValAlaGlyLeuValGlyGluAl 1013
876 GATTGACACAGCAACCGCGTATTCGCTTCGCTTATTCGCAAGCGCGCA 925
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1013 aValGluAlaTyrArgAlaLeuAlaAspGlyLysValValAspGlyThrV 1030
926 TT.....ACACAAGCGCGCAC... 942
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943 .....GATTATTGGCAAGCTACCAACATCATTCCTCCGTTATCGAAGA 986
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1047 ProGluLysTyrIle..AsnAlaGluArgLeuArgLeuGluIleTyrArgL 1063
987 AGCCCGCAGCAAGAGCTGTTCGCTGGCTGGCGCGCAGCGCAAAAT 1036
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1063 yLeuAlaIleGlnSerGluSerGluValaAspLeuArg..LeuAlaValGluG 1079
1037 ACTCCATCAGCGGTACACCCCTGGCGCATTTCTTAAACAACTCTTC 1086
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1079 uMetGluAspArgTyrGlyProIleProGluGluValGluArgLeuLeuA 1096
1087 AAGTTCAAACAGACCGGTCAACGCGCGCGCGCCAT..... 1124
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1096 lValaSerArgLeuArgHisLeuMetArgGluAlaHisIleThrAspIle 1112
1125 GGTCCGATGTGACTTACGAGCGCGTGATGCCCTGGATATCTGCCCA 1174
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1113 AlaValGln.GlyThrArgIleLysValHisProValaAspLeuAlaAspS 1129
1175 CC.....CTGCTTTGGCGGATTTAATGTCGGCATTCGACAGCGCG 1218
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1129 eArgGlnValArgLeuLysArgLeuPheProGlyAlaThrTyrArgAla 1145
1219 CAGGCATTTGGTGTCTTGAATTTGACGAGAAGACCTCGTTGTGCAG 1268
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1318 .....CTGGAA 1323
1170 sPLeuLeuGlnTrrPValAlaAsnPheIleSerAsnMetPheAsnLeuGln 1186
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seq_name: /SIDSI/gcgsdata/geneseq/geneseqp_emb1/AA2001.DAT.AAB59827

seq_documentation_block:

ID AAB59827 standard; Protein; 1592 AA.

XX AAB59827;

XX 04-APR-2001 (first entry)

XX Protein #4 encoded by Tufd/E gene.

XX Toluene degradation; enzyme; waste degradation; TufE; TufD.

XX Thauera aromatica.

OS Xanthomonas maltophilia.

```

OS Geobacter metallireducens.
OS Azobars tolulyticus.
XX WO200072650-A2.
XX 07-DEC-2000.
XX 24-MAY-2000; 2000WO-US14298.
XX 01-JUN-1999; 99US-0323872.
XX (UYOH-) UNIV OHIO.
XX Coschigano PW;
XX WPT; 2001-041080/05.
XX DR N-PSDB; AAF23627.
XX PT Composition comprising toluene degrading enzyme useful for biological
XX treatment of organic compounds, especially for degrading toluene or its
XX analogs.
XX PS Disclosure; Fig 12; 122pp; English.
XX CC The present invention relates to toluene degrading enzyme genes and
XX proteins tufd (see AAF23629 and AAB59831), tufE (AAF23630 and AAB59832),
XX tufF (AAF23631 and AAB59833) and tufG (AAF23632 and AAB59834). The
XX toluene degrading enzymes are homologues of pyruvate formate lyase. The
XX toluene degrading enzymes are useful for biological treatment of organic
XX compounds and in particular for the degradation of toluene and its
XX analogs contained in liquid or solid waste source. The present sequence
XX is a protein sequence encoded by toluene degrading enzyme gene, Tufd/E.
XX SO Sequence 1592 AA;

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alignment_scores:
Quality: 118.00 Length: 584
Ratio: 0.599 Gaps: 34
Percent Similarity: 33.733 Percent Identity: 22.774

alignment_block:

US-09-303-518D-125/rev x AAB59827 ..

Align seg 1/1 to: AAB59827 from: 1 to: 1592

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1225 .....ATGCCCTGCCGCTGTGCGTATCGCGACGATTAATCGCGCAAA 1181
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740 lAglyAlaProGlyAlaThrCysSerArgArgProPheSerArg 756
1180 GCAGGCTGGCGAGCATTCAGAGGCAAGCGCGCTCGTAGTACCAATC 1131
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757 AspSer...AlaGly..ProArgAlaAlaArgPheArgCysArgAs 771
1130 GGCACCATGGCGCGCTCCGCCCGCTTGACGGCTGTGTTGAACCTGAGAG 1081
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771 palAcysGluArgArgAlaArg..... 778
1080 TTGTGTTTTCAGAGAAATGCGGAGGCTGTACGCTGATGAGATTTGT 1031
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779 .....Cys 779
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780 ProGlyProArgSerArgProSerIleArgArgGlySerArgAspArgSe 796

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809 ...AlaThrAlaThrSerCysProAlaArgArg... 818
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586 ...CAGCTGCTTACAACATGATTTTGCCTTCG... 554
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478 ...CCAGCGGATTCG...T 466
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944 ys...ArgArgTrpArgArgProLeuLysSerSerProArg 956
439 ...ACGGCTCGGCACTGACGCGCAGAAATTTGCTGAACGAGCGG 399
957 AlaThrCysThrAlaArg...CysGlyArgAspG 967
398 GTGCGCGAGCG... 389
967 yCysSerAlaPhePheGlyAsnProLeuHisArgSerLeuArgGlyProT 984
388 ...CAGTCCACAACCGG... 374
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374 ... 374
1001 AlaValArgGlySerSerArgHisAspArgThrAlaSerThrArgArgPr 1017

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373 .....ATTGATCAAGTTGGGCGG 355
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1034 yStrProAlaGlyThrAlaSerSerArgAlaAlaSerGlyAlaSerAlaLys 1050
307 CAACCTCGATTTGTCGCT...TGCCTTCAACGGCAATCAACGACTGAC 264
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1051 ArgThrArgLeuArgArgArgSerCysProValArgSerProArgArgPr 1067
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1067 gGlyThrArgAlaAlaThrPheHisSerAlaCysGlySerSerArg... 1082
222 GCCTGAACCGCGCGCAGTAACACCA...CGCCCGGAT 188
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1083 ...ArgProSerSerGlyArgProTrpSerValProIleArgProSer 1097
187 TCTTTTGTCTTCAACACGACTTGCCTT... 158
1098 SerIleCys...GlyArgAlaValGlyLeuThrSerProSerSerProLe 1113
157 .....TTTGACGGCATCGCCTTCTGACTTTCATTCAGAGGCGG 118
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1128 ySerArgHisAsnArgAlaArgArgTyrGlySerArgArg...Prohe 1142
67 CGTCGTAA... 59
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seq_documentation_block:
ID ABG22380 standard; Protein; 2447 AA.
XX
AC ABG22380;
XX
DT 18-FEB-2002 (first entry)
XX
DE Novel human diagnostic protein #22371.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
XX 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX

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      2206  AsnLeuSerGlyGlyIleAspArgArgGlyGlnIleLeuArgPheAs 2222
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      1137  TACT.....TACGAGCGCGTATGCTGATGATGCTGGCGCACCC 1177
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      2222  nThrArgThrGlyLeuGlyValIleProValAspIleSerThrAla 2239
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AC   AAB59817;
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XX   04-APR-2001 (first entry)
XX
XX   TultD protein #8.
XX
XX   Toluene degradation; enzyme; waste degradation; TultD.
XX
XX   Thauera aromatica.
XX   Xanthomonas maltophilia.
XX   Geobacter metallireducens.
XX   Azotarcus toluilyticus.
XX
XX   WO200072650-A2.
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XX   07-DEC-2000.
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XX   24-MAY-2000: 2000WO-US14298.
XX
XX   01-JUN-1999: 99US-0323872.
XX
XX   (UYOH-) UNIV OHIO.
XX
XX   Coschigano PW;
XX
XX   WPI; 2001-041080/05.
XX   N-PSDB: AAF23625, AAF23627.
XX
XX   Composition comprising toluene degrading enzyme useful for biological
XX   treatment of organic compounds, especially for degrading toluene or its
XX   analogs
XX
XX   Disclosure: Fig 5; 122pp; English.
XX
XX   The present invention relates to toluene degrading enzyme genes and
XX   proteins tult (see AAF23629 and AAB59831), tult (AAF23630 and AAB59832),
XX   tult (AAF23631 and AAB59833) and tult (AAF23632 and AAB59834). The
XX   toluene degrading enzymes are homologues of pyruvate formate lyase. The
XX   toluene degrading enzymes are useful for biological treatment of organic
XX   compounds and in particular for the degradation of toluene and its
XX   analogs contained in liquid or solid waste source. The present sequence
XX   is a protein sequence for toluene degrading enzyme, TultD.
XX
XX   Sequence 999 AA:

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alignment_scores:
  Quality: 117.00      Length: 570
  Ratio: 0.582        Gaps: 33
  Percent Similarity: 35.263  Percent Identity: 23.158

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alignment_block:
US-09-303-518D-125/rev x AAB59817

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112  CysThrSerArgGlyArgSerArgCysSerProAspArgCysArgArg 128
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1273  .CGAAGCTGCACAAAGCGAGTCTTTGCTGCCAATTCACCAACGCA 1226
|||:|||||
128  pSerArgCysSerSerProSerProArgArgCysProProSerSerPro 145
|||:|||||
1225  ....ATGCGTGGCGGCTGCGGTATCGCGGAGCAATTAATGCGGCA 1181
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145  IagIyAlaProGlyAlaThrCysSerArgArgProPheSerArgSer 161
|||:|||||
1180  GCAGGTGGCGAGATATCCAAAGGCGATCAGCGCTGTAACTACCAATC 1131
|||:|||||
162  AspSer...AlaGly...ProArgAlaAlaArgPheArgArgCysArg 176
|||:|||||
1130  GGCACCATGGCGGCGGCGGCGGCTGACGCGCTGTGTAAGTGAAGAG 1081
|||:|||||
176  pAlaCysGluArgArgAlaArg..... 183
|||:|||||
1080  TTTGTTTTCAGGAAATGCGCGGAGGTTGTACGCTGATGAGTATTGT 1031
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184  ..CysProGlyProArgSer.....Ala 190
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1030  CCGGCTGCGGCGGCGGCGGAGCGGAGCGGCGGCTGCGGCTGCG 981
|||:|||||
191  ProSerIleArgArgGlySerArgAspArgSerArgSerArgSerArg 207
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980  ATACGCAATCTGAT...TGGTAGGCGTCCCAATATATGCGGCGCC 934
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207  gSerArgGlySerProLeuCysGlyAlaThrAlaThrSerArgProArg 223
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933  TTGTGTAATCGCGGCTTCATATCCGAACCGGAATACGCGGTGTCTG 884
|||:|||||
224  ....ArgAlaArgCysSerIleGlyAlaSerSerGlyCysPro 236
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883  TGTCAACCAATTCGCGCGGAGATTTGCGATCTTGTGCAACCCAAACG 834
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237  HisProProValArg..... 241
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833  GTACGCAAGAGCGGCGGCTTGTGACTGTAGACACCACTAGGCAATAC 784
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242  ....ArgSerProVal.....AsnSerSerIysArgAla 252
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783  GCGCTCGGCTTTCAGACGCGCTGTTCAAACAAAGGCGCAATGTTA 734
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252  IsArgArgCysThr..... 256
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733  CATCTGATTAATGATGTCACACGCGTTTATTCGCGCGGCGGCTCG 684
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257  ....AlaArgArgGlyArgPheArgGlyPro..... 265
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683  ATGAATGATATGCGTGCACCTCAACCGGCGAGATGCGGCGCGGCA 634
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266  ....ThrSerArgAspPheCylArgArg. 273
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633  TTCAATGTTCGATGTTGCGACGATTTTCAAGCGACGCTGCGC... 587
|||:|||||
274  ....ArgCysThrArgGlyProArgProArgArgCysArgCys 286
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586  ....CAGTCCCTTACAAACATGATTTTTCGCTGCG..... 554
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287  SerArgArgGlyArgProLeuThrAlaSerArgCysProArgAla 303
|||:|||||
553  ....TCAACGCGCTCATATACCAACAGCGCGGCTTGAATC 517
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303  gTTPArgArgGlySerAsnThrPheSerArgGlyArgSerSerAlaSer 320
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516  CTCGCGGCGCTTCTTGTGATTAATGACCGTAGGCTGCGCAG..... 479
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320 rQlySArgThrcys.....GLYArgValArgSerAsp 331
478 .....CCAGCGATTGG.. 467
332 ThSerAlaArgSerArgCysProAlaSerProIleArgTrpH 348
466 .....TGTCATCGCATTGACGAGATGGCGA..... 440
348 rGlyArgCys.....ArgArgTrpArgArgProLeuGlyCys 361
439 .....ACGCTCGCATCGACGCGAGGATTTGGTGA 407
361 erProAlaIaThrcysThrAlaArg.....CysGly 371
406 ACGACGGGTCGCGACG..... 389
372 ArgAspGlyCysSerAlaPhePheGlyAsnProLeuHisArgSerLeuAr 388
368 .....CAGTCCACAAACC 376
388 gGlyProTrpAlaAlaProPheArgAlaHisArgSerArgSerThr 405
375 GG..... 374
405 rGArgCysAlaValArgGlySerSerArgHisAspArgThrAlaSerThr 421
373 .....ATTGATCAGG 363
422 ArgArgProHisLysProProLysGlyCysAlaThrAspIleHisSerG 438
362 TTGCGCGCGCATTCTCCGCGCTTAAGTTTCCAGCGCTTCAAGTGGCTA 313
438 YArgTyrCysTrpProArgThrAlaSerSerArgAlaAlaSerGlyAla 455
312 GGGT...CAACTCGATTTCGCT...TGCTTCAACGGCAATCA 272
455 erAlaLysArgThrArgLeuArgArgSerCysProValArgSerPro 471
271 CGACTGACTGAAGTACGCGCTTTGCGCAC.....GGTGAATCGCG 231
472 ArgArgArgGlyThrArgAlaIaIaTrpHisSerAlaCysGlySerSer 488
230 GGGATTTCCTCCGAGCGCGCGCAGTAACACCA.....C 196
488 rArg.....ArgProSerSerGlyArgProTrpSerValProIleA 502
195 GCCCGGATTCTTTGCTTCAACAGACGACTTGGCCTT..... 158
502 rGProSerSerIleCys...GLYArgAlaValGlyLeuThrSerProSer 517
157 .....TTTGAGCGGCAATCGCTTCCCTTGACTTTCATC 126
518 SerProLeuAsnArgProPheAlaArgArgSerAlaPro.....AlaSe 532
125 GAGGCGCGCATACCGCATATCTTCGCCAAGACAGCGACTTCGGTAAT 76
532 rThrProCysArgArgHisAsnArgArgArgGlyGlySerAlaArg... 547
75 GCGCGGCGCGTCGTAA..... 59
548 ...ProPheArgArgArgPheAlaCysSerTrpSerSerGlnHisAspPro 563
58 .....CGGCTTGCCTCGCGTTCGCGC 39
564 AlaSerGlnAspProGlnArgGlyThrCysProLeuArgAsnAlaCysPr 580
38 GCGATGGGCA 29
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seq_name: /SID1/gcdata/geneseq/genesep-emb1/AA27242

seq_documentation_block:

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ID AAB27242 standard; Protein: 571 AA.
XX AC AAB27242;
XX AC 27-MAR-2001 (first entry)
XX DT 27-MAR-2001 (first entry)
XX DE Human EXMAD-20 SEQ ID NO: 20.
XX OS Homo sapiens.
XX PN WO20068380-A2.
XX PD 16-NOV-2000.
XX PF 10-MAY-2000; 2000WO-US12811.
XX PR 11-MAY-1999; 99US-013643.
XX PR 23-AUG-1999; 99US-0150409.
XX PA (INCYTE) INCYTE GENOMICS INC.
XX PI Bandman O, Hillman JL, Tang YT, Lal P, Yue H, Baughn MR, Lu DM;
XX PI Azimzai Y;
XX DR WPI: 2001-007395/01.
XX DR N-PSDB; AAC66909.
XX PT Isolated polynucleotide encoding extracellular matrix or
XX PT adhesion-associated protein (EXMAD) useful for diagnosing, treating, or
XX PT preventing disorders associated with expression of EXMAD such as
XX PT proliferative, immune and genetic disorders -
XX PS Claim 1: Page 106-107; 129pp; English.
XX CC The present invention provides the protein and coding sequences for 25
XX CC novel extracellular matrix and adhesion-associated proteins (EXMADS).
XX CC These are designated EXMAD-1, EXMAD-2, EXMAD-3, EXMAD-4, EXMAD-5,
XX CC EXMAD-6, EXMAD-7, EXMAD-8, EXMAD-9, EXMAD-10, EXMAD-11, EXMAD-12,
XX CC EXMAD-13, EXMAD-14, EXMAD-15, EXMAD-16, EXMAD-17, EXMAD-18, EXMAD-19,
XX CC EXMAD-20, EXMAD-21, EXMAD-22, EXMAD-23, EXMAD-24 and EXMAD-25. They are
XX CC useful in the prevention and treatment of cancers, cell proliferation,
XX CC cardiovascular, reproductive, immune, musculoskeletal, developmental and
XX CC gastrointestinal disorders and inflammation.
XX SO Sequence 571 AA;

```

alignment_scores: Quality: 116.00 Length: 472
Ratio: 0.560 Gaps: 20
Percent Similarity: 43.856 Percent Identity: 19.703

alignment_block: US-09-303-518D-125/rev x AAB27242 ..

Align seg 1/1 to: AAB27242 from: 1 to: 571

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1290 GTATTGCCGCGGAGACGAGACGCTGACAAAGCGAGCTCTTCGTCGA 1241
39 nThrAlaLysThrSerThrSerLeuHis.....SerHisThrSers 53
1240 ATTCCAAAGCAACCAATGCTGCGCGGCTGTCGGTA..... 1206
53 erThrHisHisProGlnValThrProThrSerIleThrAsnIleThrLeu 69

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86 ThrSerSerAlaLeuThrThrProThrThrHisSerProThrG 103
1129 GCACC.....ATGCGCGGTGCGCGCTGACGGCTGTGTAACCTG 1086
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1035 TTGTCGCGGTGCGCGCAACCGCAACAGCTTTGTGCGGCTT 986
127 .....ProSerHisProGlnThrThrLeu..... 134
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818 GGTTTGTGACTTGAACCACTAGGCAATC..... 786
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785 .....ACGCGCTGCTTTCAGACGCGCTTTCGAAACAAACG..... 747
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201 rGlnThrHisSerSerPheSerThrAlaThrAlaSerSerPheIles 218
746 .....CCAAATGTAATTAATCATCTTGATTA 723
218 eIleSerSerSerThrSerSerThrPleuProGlnAsnSerSer..... 232
722 TTGATGTCCACAGGTTTATTCGCGCCGACCGCTCGATGAATGAAT 673
233 .....ArgProProSerSerProIleThrTh 241
672 GTGCGTCCACTCAAAACCGGAGATGCGGCGCAATTCATGTGTTT 623
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241 rGlnLeuProHisLeuSerSerAlaThrThrProValSerThrThAsnG 258
622 CGATGTGGACGATTTTTCAGCGGACGCTGCGCACCTGCTTACAA 573
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258 InLeuSerSerPheSerProSerProSerAlaProSerThrValSer 274
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522 GAATCTCGCGCGCTTCTTGTATATGACCGTAGGCTGCGGACGACG 473
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291 IglThrSerSerSerPheValSerAlaProValHisSer..... 304
472 GATTGGTTCATCGCATTCGACGAAGATGCGACGCGCTCGGCATCG... 426
305 .....ThrThrLeuSerSerGlySerHisSerSer 314

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425 .....ACGCGACGAAATTTGCTGACAGCAGCGGTGCG 394
315 LeuSerThrHisProThrThrAlaSerVal..... 324
393 CACGCGAGTCCACAAACCGGATGTGATCAGGTTGCGCGCACTGTGCG 344
325 .....SerAlaSerP 328
343 CGCTTAAGTTTGGCAGCGCTTACGTTGCGTACGTTCAAACTGAT... 297
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296 TCGTCGTGCTTTCACAGCGCAATCAACGACTGACTGAATACCGCTTTC 247
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199 CCACGCGCGGATTTCTTTGTCTTCAACAGCAGCTTGGCTTTTGACG 150
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seq_name: /SID1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:ABG03530

seq_documentation_block:

ID ABG03530 standard; Protein; 599 AA.

XX ABG03530;

DT 13-FEB-2002 (first entry)

DE Novel human diagnostic protein #3521.

KW Human; chromosome mapping; gene mapping; gene therapy; forensic;

KW food supplement; medical imaging; diagnostic; genetic disorder.

OS Homo sapiens.

XX W0200175067-A2.

XX 11-OCT-2001.

PF 30-MAR-2001; 2001MO-US08631.

XX 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

XX (HYSE-) HYSEQ INC.

PI Drymanac RT, Liu C, Tang YT;

XX WPI; 2001-639362/73.

DR N-PSDB; AAS67717.

PT New isolated polynucleotide and encoded polypeptides, useful in

PT diagnostics, forensics, gene mapping, identification of mutations

PT responsible for genetic disorders or other traits and to assess

CC Claim 20; SEQ ID No 33889; 103pp; English.

CC The invention relates to isolated polynucleotide (I) and

CC polypeptide (II) sequences. (II) is useful as hybridisation probes,

CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II) (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. ABG0010-ABG30377 represent novel human
CC diagnostic amino acid sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_prt_sequences.

XX Sequence 599 AA;

alignment_scores: Quality: 116.00 Length: 520
 Ratio: 0.563 Gaps: 36
Percent Similarity: 39.615 Percent Identity: 26.538

alignment_block:
US-09-303-518D-125 x ABG03530 ..

Align seg 1/1 to: ABG03530 from: 1 to: 599

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38 o...ArgTthAlaAlaGly***ArgArgTyrTyrCysAlaSerAlaSerL 54
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124 TCGATGAAGTCAGAGAGCGATGCGCTGCAAAAAGGCCAAGTCTGTT 173
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54 euAla**SerPro.....CysArgProArg...SerArgTyr 66
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174 TGAAGCAAAAAGATCCGGCGCTGCTTACTGC..GCCGGCTTCAG 220
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67 TTPArgAspAlaGlySerGlyTyrThrProHisCysProAlaSerAla 83
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265 TCAAGTCGTATGCCGTTGAAGCAACGACGAATCGATTGAACGCTA 314
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100 InSerHis..... 102
315 CGCACCTGAAGCGCTGGCAAACTTAAGCGGCAAGAAGTGGCGGCAAC 364
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103 .....CysProGlyGlyLeuArg.....AlaProProPr 112
   |||||
365 TGATCATTCGCGGTTTGACTGCGC.....TGCGACCGCTGCTCAG 408
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112 oglySerValArg..CysSerThrGln***AspCysSerSerValArgPr 128
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409 AGCA.....AAATCTCTCCGCTGCATGC 431
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128 oAlaTyrSerArgSer***GlyAlaCys***GlnVal**ProArgCysP 145
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432 CGAGCGTGGCCATCTTCGCAATGCGATGACACCAATCCGCTGCTG 481
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145 rOCysArgThrProAlaThrLeu.....TrrAlaPro..... 155
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617 ACAT.....C 621
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197 ahisProGlyAlaLysSerLeuGlyLeuAlaCysGlnProHisArgGlyL 214
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622 GAACACATGAATTTGGCGGCGCGCATCTCGCGGTTTGAAGTGGACGCA 671
   |||||
214 ysglyThrProLleGlnGlyProAla...CysGlyThr***GlyGlyArg 229
   |||||
672 CATTCATTTTCATCGAGCGCGGTCGGCGGCAATMAAACGCTGGACATCA 721
   |||||
230 ArgGlySerGlyCysProGlyArg.....ProHisTh 240
   |||||
722 ATTATCAAGATGTATTTACCATTTGGCGCTTGTTCACACAG..... 764
   |||||
240 rArgArgArgCys***Pro.....ProAlaProCysGlyArgArgSerA 255
   |||||
765 ..CCGCTGGAACACCGGCGGCTGATTCGCCGTAGGTTCTCAAGTCAA 812
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255 laGlySerAlaHisProAlaThrProThrProHisGlyProGlyGlyLn 271
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   |||||
272 GlnArgAspProGlyProAlaTyrArgGlyGlyGlnGlyAlaArgSerPr 288
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854 CGCAATTTACTGCGCGCAATTTGGTTGACACAGACAAACCGCGTATTC 903
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288 oAlaSerProSerGlyArg.....A 295
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904 GGTTCGATTTGAACGCGCGCATTTACAAAGCGCGCACGATTTTGGG 953
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295 rGlnProAlaSerArgAlaGlyArgSerArgAlaAlaArg.....Gly 309
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1170 GCC.....CACCT.....GCTTTGGCGGATTT 1192
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407 AlaArgAlaArgArgHisProGlyArgCysProGlnAlaSerGlyProAr 423
   |||||
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1224 .....ATTGGTTCCTGGAAATGAGCAAGAGACT 1256
440 InProGlyAlaProProCysHISLeuProGlyIleProAlaArgGlnPro 456
1257 CGCTTTGTCAGCTTCGTCTGCCCGGCAATAGCAATAGCGCCGCTCT 1306
457 .....LeuGlyLeuProArgArgThrArgCysPheGlyGlyI 469
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469 eAlaGln 471

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seq_documentation_block:
ID ABG03731 standard; Protein; 696 AA.
XX
AC ABG03731;
XX
DT 13-FEB-2002 (first entry)
XX
DE Novel human diagnostic protein #3722.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Dmanac RT, Liu C, Tang YN;
XX
DR WPI: 2001-639362/73.
DR N-PSDB; AAS67918.
XX
XX
XX New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity
XX
PS Claim 20; SEQ ID No 34090; 103bp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX (II). (II) is useful for generating antibodies against it, detecting or
XX quantifying a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations in
XX responsible for genetic disorders or other traits to assess biodiversity
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. ABG0010-ABG03737 represent novel human
XX diagnostic amino acid sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed

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CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 696 AA:

alignment_scores:
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  Ratio: 0.550        Gaps: 24
  Percent Similarity: 45.671  Percent Identity: 23.810

alignment_block:
US-09-303-518D-125 x ABG03731 ..

Align seg 1/1 to: ABG03731 from: 1 to: 696

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59  TTACAGCGCGCCGCCATTACGAGTCCGCTTGGCCAGAAATAT 108
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27  g***ArgArgValSerProThrArgSer.....GlyLysArgArgGly 42
109 GCCGTATCGCGCCCTCGATGAAGTCAAGAGAGCGGATCGCTCAAAA 158
   ::::  ::|||:::  |||  :::::  ::
42  IagIuIuLysAsnArgGlnGluLysLysGlyArgGluIuLysGluArg 58
159 AGGCCAAGTGTCTTGAAGACAAAAGAAATCCGGCGCTGTACTG 208
   ||  ::  |||||:::  |||
59  ArgGluLysArgSerGluArgGlnArgAspArgArgArg..... 72
209 CGCGCGCTTCAGGCAAAATCGCGCGATTCACCGTGGCAAAAGCGCTA 258
   :::::  ::|||:::  |||  ::|||:::  |||
73  .....LysGluGlnArgLysGluIuLgnArgArgAlaArg 86
259 CTTCAGTCAGTCGTGATTCGCTGGAAGCAAGCAAGAAATGAGTTGA 308
   ||::  ::|||:::  |||  ::|||:::  |||
86  hrAsn.....GluArgLysProArg..... 92
309 ACGCTACGACCTGAAGCGCTGGCAAACTTAAAGCGGAGAAAGTGGCC 358
   ::|||  ::  |||  ::  ::  ::|||:::
93  .....GlnThrGlnAlaAsnGlyAlaThrSerSer**LysAlaSerAl 107
359 GCACCTGATCCAAATCCGGTTGTGGACTGGCGTCCGACCCGCTTC 408
   |||  ::  ::  ::|||:::  |||:::  |||:::
107  eGlnGlnAlaGlyMet.....TyrGlyLysPro**Thr 120
409 AGCAAAATTCCTGCCGTGATGCCAGCGCTTGGCATCTTCATCATGC 458
   ::  ::::  |||||  |||
120  spAlaThrAlaIleArgArgGlyAlaProCysSerSerArgArgThr 136
459 GATGACACCAATCCGCTGGCTGCCAGCCCTTACGTCATTTCAAAGAG 508
   ::|||:::  ::  |||  |||
137  CysLeuAsnGlnGlyThrIleAlaThrProSerGly.....ArgAr 150
509 CGCGCGAGGATTTCAACGCGCGCTGTGTATTCAGCCGTTGACCGAA 558
   ||:::  |||  |||  |||:::  ::
150  gArgHisGlyAspAlaGly***Pro..Gly.LeuAlaSerGlnHisAsp 165
559 CGCAAAATTCATGTTGTAAAGCAAGTGGCGAGACAGTCCGCTGAAA 608
   ::  |||  |||  ::|||  |||
166  AlaSerGlnHisGlyCysLeuArgThrGlyAlaGly**ProSerAspSe 182
609  T.....GCTGCCAATCATGCAACACATG 631
182  ThrGluSerValLysArgArgProLeuAlaMetHisValProThrHisG 199
632  AATTCGGCGCGCCGACATCCGCGTTCGATGAGTGCAGACATCATTCG 681
   ||  |||||  ::::  |||:::  |||
199  IuSerHisGlyPro.ValPheThrArgLeuValSerHisThrPheHisCy 215
682  ATCGAGCGCGTGGCGGCAATA...AAACCG.....TGTGACCATCA 722

```


[illegible]

```
seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2000.DAT:AAAB159355
seq_documentation_block:
ID   AAB15935 standard; Protein; 740 AA.
```

DT	05-OCT-2000	(first entry)
XX		
DE	E. coli proliferation associated protein sequence SEQ ID NO:292.	
XX		
KW	Escherichia coli; E. coli; proliferation; inhibition; screening;	
KW	antimicrobial; bacterial growth; antisense therapy; antibacterial	
XX		
OS	Escherichia coli.	
XX		
PN	WO2000/44906-A2.	
XX		

PD 03-AUG-2000.
 PE 27-JAN-2000; 2000WO-US02200.
 XX
 PR 27-JAN-1999; 99US-0117405.
 XX
 PA (ELIT-) ELITRA PHARM INC.
 XX
 PI Zyskind J, Ohlsen KL, Trawick J, Forsyth RA, Froelich JM, Carr GJ,
 PI Yamamoto RT, Xu HH;
 DR WPI; 2000-514822/46.
 DR N-PSDB; AAA65940.
 XX
 XX Novel polynucleotides and polypeptides associated with microorganism
 PT proliferation, used to identify inhibitors of bacterial growth and
 PT proliferation, for use in antisense therapy -
 XX
 PS Claim 11; page 217-219; 316pp; English.

AA
CC AAA65809 to AAA65889 and AAA66058 to AAA66138 represent nucleotide
CC sequences derived from *Escherichia coli* which inhibit *E. coli*
CC proliferation. AAA65890 to AAA66055 and AAB15886 to AAB16040 represent
CC nucleotide and protein sequences associated with *E. coli* proliferation.
CC AAA66056 and AAA66057 represent primers used for sequencing *E. coli*
CC proliferation inhibiting nucleotide inserts in an example from the
CC present invention. Methods from the present invention can be used to
CC identify a proliferation- required gene in a microorganism, by contacting
CC a microorganism with a proliferation-regulated gene actively inhibitory
CC to nucleic acid identified in another organism, and determining if
CC inhibition occurs in the second microorganism. The nucleic acid sequences
CC identified as being required for bacterial growth and proliferation, can
CC be used for antisense therapy for killing bacteria.

Sequence 740 AA;

```
alignment_scores:
  Quality: 115.00      Length: 322
  Ratio: 0.728        Gaps: 12
Percent Similarity: 49.068  Percent Identity: 22.050
```

```
alignment_block:
US-09-303-518D-125 x AAB15935 ..
```

Align seg 1/1 to: AAB15935 from: 1 to: 740

```

100 GAAAGATATTCGGCTATGGCGCCCTGATGAAAGTCAAGGAAGCGCATGC 149
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
48 LysGlnHisIleGlyIaGluGlyLeuGlyValSerValGlyAspLys 64
150 CGTCAAAAAGCGCATGCTGTGTTGAAGACCAAAAAGCATCGCGCGCTGG 199
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
64 sValLeuATGtGlyGlnProLeuThraArgIYArgGlyLysMetLeuProV 81
200 TGTATTACTGCGCGCGCTTCAGGCAAAATCGCCGCGATT..... 237
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
81 aHis...AlaProIlnSerGlyThValThAlaIleAlaIleProHisSer 96
238 .....CACCGTGGCGAAAAGCGCGCTACTTCAGTCAGTCGATTGCCGT 281
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
97 ThrAlaHisProSerAlaLeuAlaGluLeu...SerValIleIleAspAl 112
282 TGAAGCAACGAC..... 294
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
112 aspGlyGlnAspCysTrpIleProArgAspGlyTrpAlaAspTyrArgT 129
295 .....GAATTCGAGTTTGAAACGCTAGCAACCTGGAAGCGCTGGCA 333
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
129 hTrArgSerArgGlnGluLeuIleGlnArgIleHisGlnProGlyValAla 145
334 AACTTAAGCGCGCAAGAAAGTGGCGCCCAACGTCGATTCGAAATCGGTTGTG 383
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

```

146 GlyLeuGlyGly.....AlaGlyPhePr 153
 384 GACTGGCTGCGCACCCGCTCCGTCAGCAAAATTCCTGGCGGATGGCG 433
 153 oThrGlyVal.....LysLeuGlnGlyGlyAspL 164
 434 ACCCGTGGCCATCTTCGTAATGCGATGACACCAATCCGCTGGCTGCC 483
 164 yAlaGlnThrLeuIleIleAsnAlaIleGluGlyGluProTyrIleThr 180
 484 GACCCACGCGCATATATCAAGAACGCCGCCGAGATTCCAACGCGGCT 533
 181 AlaAspAspArgLeuMetGlnAspCysAlaIleGlnValIleGluGly 197
 534 GTTGGTATGAGCCGTTGACCGAA..... 558
 197 eArgIleLeuAlaHisIleLeuGlnProArgGluIleLeuIleGlyLeu 214
 559CGCAAAATCCATGTTGTAAAGCAGCTGGCGCAGC 594
 214 LuAspAsnLysProGlnAlaIleSerMetLeuArgAlaValLeuAlaAsp 230
 595 GTGCGCTGAAAAATGCTGCCAATCGAACACATGATGATTCGGCGGCC 644
 231SerAsnAspIleSerLeuArgValIleProThrLy 242
 645 GCATCTGCGCGGT.....TTGAGTGGCAGC 670
 242 sTyrProSerIleGlyAlaLysGlnLeuThrTyrIleLeuThrIleGly 259
 671 ACATTCATTTATGAGCCGCTGCGCGCAATTAACCGCTGGACCATC 720
 259 LuVal.....ProHisGlyArgSerSerAspIleGlyVal 271
 721 AATTATCAAGATGTAATTACC.....ATTGCGCGTTGTTGCAAC 761
 272 LeuMetGlnAsnValIleGlyThrAlaTyrAlaValLysArgAlaValIleAs 288
 762 AGCGCGCTGTAGACACGAGCGCGTATGCCCTAGTGTGTTCCAACTCA 811
 288 pGlyGluProIleThrGluArgValIleThrLeuThrGlyAlaIleAs 305
 812 ACAACCGCGCGCTCTGCTACCGTTTGGTGGGAAGATGCAAAAT 861
 305 IaArgProGlyAsnValIleThrAlaArgLeuGlyIleProValArgHisLeu 321
 862ACTGGGCGCAATTGGTTGACACAGAACCGCGTATTCGG 905
 322 LeuAsnAspAlaGlyPheCysProSerAlaAspGlnMetValIleMetG 338
 906 TTCGATTTGAGCGC 921
 338 yGlyProLeuMetGly 343

seq_name: /SIDS1/gcgdata/geneseq/geneseq-p-emb1/AA1999.DAT:AAV04955

seq_documentation_block:

ID AAV04955 standard: Protein: 573 AA.

AC AAV04955;

DT 06-JUL-1999 (first entry)

DE Mycobacterium species protein sequence 41T#3.

KM Secreted protein: Mycobacterium; primer: PCR; amplification: probe;

KW hybridisation: detection; vaccine; immunisation; infection.

OS Mycobacterium sp.

XX W09090186-A2.

XX 25-FEB-1999.

XX
 PF 14-AUG-1998; 98MO-FR01813.
 XX
 PR 11-SEP-1997; 97ER-0011325.
 PR 14-AUG-1997; 97ER-0010404.
 XX
 PA (INSP) INST PASTEUR.
 XX
 PI Gicquel B, Lim EM, Pellicle V, Portnoi D, Goguet de la Salmoniere Y;
 PI Guineno A;
 DR WPI: 1999-181045/15.
 DR N-PSDB: AAX34206.
 XX
 PT Mycobacterial DNA vectors containing reporter constructs - for
 PT Identifying coding or promoter sequences involved in
 PT infection-associated protein expression
 XX
 PS Claim 32; Fig 41T; 309pp; French.
 XX
 CC Sequences AAV04742-Y05000 and AAV07201-Y07204 represent secreted
 CC proteins from various Mycobacterium species microorganisms. The
 CC encoding nucleotide sequences can be used as primers and probes for
 CC methods for detecting and identifying mycobacteria, especially belonging
 CC to the M. tuberculosis complex. The encoded proteins can be used in
 CC vaccines for immunisation against a bacterial or viral infection.
 XX
 SO Sequence 573 AA;

Alignment_scores:
 Quality: 114.00 Length: 472
 Ratio: 0.576 Gaps: 24
 Percent Similarity: 41.949 Percent Identity: 25.424

Alignment_block:

US-09-303-518d-125 x AAV04955 ..

Align seg 1/1 to: AAV04955 from: 1 to: 573

36 CCGCGGACACCGGAGCAAGCCGTTACGACGCCCGCCATTCACGAG 85
 162 ArgGlyGlyAlaGlyAsnTyrIleGlnGlyAlaIleGlyArgArgSe 178
 86 TCGC.....GTTGCTGGCGAAGATATGCGCGATCCGCTCGATG 129
 178 rArgArgProValArgAlaArgGlyValGlyArgCysGlyHisArgArg 195
 130 AAGTCAGAGGAGCGGATGCGTCAAAAAGGCCAAGTGTGTTGAGA 179
 195 rg**ArgGlyGlyHisArgAlaGlyLysAspProArgThrAla**Arg 211
 180 CAAAAGAAATCCGGCGCGTGTACTGCGCGCTTCAGGCAAAATCG 229
 212 AlaArgArgCysGlyArgGly.....GlyArgArg 221
 230 CCGCATTCACCGTGGCGAAGCGCGTACTTCAGTCAAGTCGATTGCC 279
 221 gArgThrGlyProAlaGlySerAlaGlyArgValAlaLeuHisHisLeu 238
 280 GTTGAAGCAACGACGAATGAGTTGAAACGTCACGACCTGAGGCGT 329
 238 rGlyAlaGlyThrCysProGlyGlyLeuArgGlyThrLeu.....Ala 250
 330 GCAAACTTAAGCGCGAGAGAGTGGCGGACCTGATCAATCCGCT 379
 251 AlaArgValAlaAspArgHisGlyTyrProThrProArgProAlaIle 267
 380 TGTGGAC.....TGGCTGCGCACCGCTCGC 405
 267 gGlyAspValSerValGlyGlyMet**CysCysSerGlyGlyProVal 284
 406 TTCAGCAAAATTCCTGCCGTCGATGCCGAGCGTTCCGCAATCTTCGCA 455

```

284 1a.....GlySerThrGlnGlyLeuGly***ValGlyGlyHisArgArg 298
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
456 TGGCATGGACACCAATCCGCTGGCTGGCCGACCTACGCTCATATCAAG 505
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
299 CysSerAlaArgGlnLeuLeuArgThrArgProHis...ArgArgArgAr 314
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
506 AAGCGCGCGAGATTCTCAA.....ACGCGCG 531
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
314 gCysArgArgGlySerArg11LeuGlyGlyAlaSer***ProAspArg 331
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
532 CTGTTGGATTGAGCCGTTTGACGACGCAAAATCCATGTTTGTAAAGC 581
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
331 sPLeuGlyAla.....ArgPheArgAspGlnArg11Leu1LeuGly 343
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
582 AGCTGG.....CGCAGACGTCGCGCTGAAAAATGCTGCCAACCA 619
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
344 ArgTrpLeuAspAlaGlyProArgArgAlaGly..... 354
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
620 TGGAAACACATGATTTCCGCGCGCGCATCCGCGGTTTGAGTGCACG 669
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
355 .....GlyArgArgArgArgArgCysArg..... 362
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
670 CACATTCATTTGATCGAGCCGCTGGCGCGAATAAAACGCTGTGACCAT 719
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
363 ..ArgAlaValAlaArgArgGlyGlyArg..... 370
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
720 CAATTATCAAGATGTAATTACCATTTGGCGGTTGTTGCACAGCGCGTC 769
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
371 .....LeuArgAlaAla11th, 375
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
770 TGAACACGAGCGCGGATTTGCCCTAGTGTTCTCAAGTCAACAACCG 819
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
375 rGlySerArgArgArgAspThrGlyArg..... 384
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
820 CGCCTCTTCGATACCGTTTGGGTGGCGAAAGTACGCAATTACTGCGGG 869
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
385 .....ArgTyrGlnCysProProAlaGly.....AlaGly 394
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
870 CGAATTGGTTGACACAGACACACCGCTGATTTCCGCTTGGTATTGACG 919
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
395 Arg...GlyArgHisArg.....ArgArgAlaArgAspGlyAlaAla 408
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
920 GCCCGATTACACAGCGCGGACGATTTATTTGGAGCTACACGACATGAG 969
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
408 nTrpLeuCysGlyArgArg...ArgThrGlyGlyArgVal11TrpArgGly 424
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
970 ATTTCGGTTATCGAAGAGGCGCGACGCAAAAGCTTTGCGCTGGCTGTC 1019
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
424 sp...ArgLeuGlyArgArgArgGlyThrArgAlaAspArg11Lea 439
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
1020 GCGCGACCGCGACAAATACCTCCATCGCGGTACACACCTCGGCAATTTCC 1069
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
440 AlaGlyValGlyArg.....AlaGlyArgAla***ArgGlyLyr 452
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
1070 TGAATAACAATCTTCAAGTTCAACACAGCCTCAACGCGCG..... 1112
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
452 oProGlyArgArgArgLeuGlnHisGlyProCysArgArgCysArgPhe 469
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
1113 .....CGACCGCGCGCATGCTGGCGATTGG... 1136
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
469 roAlaArg11LeuAlaHisCysHisProLys...GlyAlaAspLeuGly 484
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
1137 ...TACTTAGAGAGCGCTGATGCCCTTGATATCCGCGCGACCGCTGTT 1183
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
485 ArgTyrLeuGln11AlaGlyArgArgSerGlyTyrArgGlyAlaArg 501
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
1184 TGGCGCATTTAATCGT...CGCGCATACCGACGCGCGCGCATTTGGGT 1230
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
501 aaSPArgArgArgArgCysArgArgGlyGlyHisArgSerGlyArgPro 518
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
1231 TGCTTGAATTGACAGAAAGACCTTCGCTTGTGCAAGCTTGGT..... 1274
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::

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518 a1ValGly11LeuGlyArgArgSerGlyAspGlyAlaAsnTrpArgArg 534
1275 .....CTGCCCGCGCAAAATACGAATACGCGCGCGCTG 1306
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
535 AsnArgArgArgGlyCysArgArgProGlyThrAlaCysAlaArgPro 551
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
1307 TGGCAAAAGTCTGGA 1322
      ::      ::      ::      ::      ::      ::      ::      ::      ::      ::
551 rArgHisArgAlaGly 556

```

seq_name: /SIDS1/gcgdata/geneseq/geneseq_emb1/AA2001.DAT.AAB96063

seq_documentation_block:

ID AAB96063 standard; Protein; 822 AA.

AC AAB96063;

DT 29-OCT-2001 (first entry)

DE Putative P. abyssi pyruvate kinase #1.

KW Hyperthermophilic archaeon; hyperthermophilic protein.

XX Pyrococcus abyssi.

OS FR2792651-A1.

PN 27-OCT-2000.

PE 21-APR-1999; 99PR-0005034.

PR 21-APR-1999; 99PR-0005034.

PA (CNRS) CNRS CENT NAT RECH SCI.

PI (IFRE-) IFREMER INST FR RECH EXPL MER.

PI Querellou J, Weissenbach J, Saurin W, Hellig R;

DR WPI; 2001-126236/14.

PT New nucleotide sequences isolated from Pyrococcus abyssi encode

PS proteins useful in industry -

PS Claim 7; Pages 677-680; 1657Pp; French.

CC The present invention relates to the genomic sequence of Pyrococcus

CC abyssi (see AAF86431 and AAH41223-7) and P. abyssi proteins. P. abyssi is

CC a hyperthermophilic archaeon, which is isolated from deep sea

CC hydrothermal vents. The present sequence is one such P. abyssi protein.

CC The proteins of the present invention have various potential industrial

CC uses, since the proteins are stable at very high temperatures, some up to

CC 110 degrees centigrade.

CC Note: This patent is in the same patent family as WO200065062, which

CC contains additional sequences as shown in AAB99132-AAB99143,

CC AAH75903-AAH75920 and AAG66436.

SO Sequence 822 AA;

alignment_scores: Quality: 113.50 Length: 455

Ratio: 0.473 Gaps: 21

Percent Similarity: 52.747 Percent Identity: 21.538

alignment_block: US-09-303-518D-125 x AAB96063 ..

Align seg 1/1 to: AAB96063 from: 1 to: 822

67 GGCCTGGCATACCGAAGTCCGCTTGGCGCAAGATATGCCGGTAT 116

255 G1yG1uAlaValSerG1yAlaValThrProAspG1yTr1LeuAlG1 271

XX
DR WPI: 1997-111454/11.
XX New immunogenic fragments of Bordetella adenyl cyclase haemolysin -
PT and related nucleic acid and antibodies, for use in vaccines and
PT immunotherapy
XX
PS Claim 6; Page -: 55pp; French.
XX
CC This is the sequence of a novel mutant, designated delta-Cla, derived
CC from the Bordetella pertussis adenyl cyclase-haemolysin (AC-Hly) in which
CC amino acids 827-887 of the wild type sequence have been deleted by
CC genetic engineering. The novel protein is able to induce production
CC of protective antibodies against the Bordetella species pertussis,
CC parapertussis and/or bronchiseptica, especially in protective vaccines
CC for human or veterinary use.
CC Note: this sequence is not given in the specification but is generated
CC from the wild type sequence.
XX
SQ Sequence 1645 AA:

alignment_scores:
 Quality: 109.50 Length: 457
 Ratio: 0.537 Gaps: 26
Percent Similarity: 44.639 Percent Identity: 22.101

alignment_block:
US-09-303-518D-125 x AAW13502 ..

Align seg 1/1 to: AAW13502 from: 1 to: 1645

190 CCGGGGCTGTGTTACTGCGCGGCTTACGCAAAATCGCGCGCATTTCA 239
 ||||| ||||| ||||| ||||| |||||
694 ProvalaValValValThrSerLeuThrGlyAlaLeuAsnGlyIleLe 710
240 CCGTGGCGAAAGCGCGTACTTCACTGATCGTGGATTCGCGTTGAAGCA 289
 ||||| ||||| ||||| ||||| |||||
710 uatgglly.....ValGlnGlnProIleIleGlyIuysleuAlaA 723
290 ACGAC.....GAAATCGAGTTT 306
 ||||| ||||| ||||| ||||| |||||
723 snaspyrAlaarglyIleaspluLeuglyGlyProGlnAlaIyrPhe 739
307 GAAGCTAGCGACCT.....GAAGCGTGGCAACTTAAGCGCGCA 347
 ||||| ||||| ||||| ||||| |||||
740 GlnuysAsnleuGlnAlaargHisGluGlnleuAlaAsnSeraspely.. 755
348 AGAAGTGGCGCGCAACCTG.....ATCCATTCGGTTGTGGACTGCG. 390
 ||||| ||||| ||||| ||||| |||||
756LeuarglysmetleuAlaAspleuGlnIaagly..TyrAsnAlaAs 770
391CTGGCGACCCGTCCTGTCAGCAAA..... 414
 ||||| ||||| ||||| ||||| |||||
770 erSerValIleGlyValGlnThrThrGlnIleSerIysSerAlaLeuGln 786
415ATTCTGCGCGTGCATGCCGA 434
 ||||| ||||| ||||| ||||| |||||
787 LeuAlaAlaIleThrGlyAsnAlaAspAsnleuLysSerValasp..... 801
435 GCGCTTCGCATCTTGCATCAATCG..ATGACACCAATCGCTGCGTG 481
 ||||| ||||| ||||| ||||| |||||
802ValPheValaspArgPheValGlnGlyIuargValAlaAg 815
482 CCGACCTAGCGTCATTAATCAAGAACCGCGCGATTTCAACCGCGG. 531
 ||||| ||||| ||||| ||||| |||||
815 lylGlnProValValleuAspValAlaAlaGlyIylaspLeuAlaLys... 830
532 CTGTGTGATTTAGCGCTTGACCGAAGC..... 561
 ||||| ||||| ||||| ||||| |||||
831ValValSerGlnleuValaAspAlaAsnGlyValleuLysHis 845
562AAATCCATGTTTGTAAAGCAGCTGGCGAGACGTCGCTGGA 607

845 rIleLysLeuAspValIleGlyIylaspGlyAspAspValValleuAlaA 862
608 ATGCTGCCACATCGAAACACATGATTCGGCGCGCGCGCTGCGCGT 657
 ||||| ||||| ||||| ||||| |||||
862 snAlaSerArgIle.....HisTyrAspGly 870
658 TTGAGTGCACGACCATTCATTCATGACCGCGTGGCGCATTAAC 707
 ||||| ||||| ||||| ||||| |||||
871 GlyAlaGlyThrAsn..... 875
708 CGTGTGCACATCAATAT.....CAAGATGTATATCA 742
 ||||| ||||| ||||| ||||| |||||
876ThrValSerTyrAlaAlaLeuGlyIylargGlnAspSerIleThrV 890
743 TTGGCGCTTTGTTGCCAACAGGC...CGTGTGAACCGCGCGTGTAT 789
 ||||| ||||| ||||| ||||| |||||
890 alSer.....AlaAspGlyIuArgPheAsnValArgLys..... 901
790 GCCCTAGTGTGTTCTCAAGTCACACAAACCGCGCTTTCGT..... 831
 ||||| ||||| ||||| ||||| |||||
902GlnleuAsnAlaAsnValTyrArgGlyIyl 913
832ACGCTTTGGTGGCGAAAGTATGCCAAT..... 861
 ||||| ||||| ||||| ||||| |||||
913 lAlaThrGlnThrThrAlaTyrGlyIylargThrGlnAspValGlnTyr 930
862ACTGGCGCGCATTTGTTGCACACAGCAAC... 891
930 rGHisValGlnleuAlaArgValGlyIylValGlnValAlaAspThrLeu 946
892CGCTGATTTCCGTTCCGTTATGAAACGCGCGATTAACA 932
 ||||| ||||| ||||| ||||| |||||
947 GlnHisValGlnHisIleIleGlyIylAlaGlyAsnSperIleThrGl 963
933 AGCGCGCGACGATTATTG..... 951
 ||||| ||||| ||||| ||||| |||||
963 yAsnAlaHisAspAsnPheLeuAlaGlyIylSerGlyAspAspArgLeuA 980
952 ..GGACGCTACCAACATCAGATTCCGTTTTCGAGAAGCGCGC..... 993
980 spGlyIylAlaGlyAsnSperThrLeuValGlyIylGlnGlnAsnThr 996
993 993
997 ValIleGlyIylAlaGlyAspAspValPheLeuGlnAspleuGlyValTr 1013
994 .AGCAAAAGCTGTTCGGCTGGTGGCGCGCGCGCGCAACATATCA 1042
 ||||| ||||| ||||| ||||| |||||
1013 pSerAsnGlnleuAspGlyIylAlaGlyValAspThrValIylsTyrAsnV 1030
1043 TCACGCGTACAAACCTCGGCCATTTCGAAAAACAACTCTTCAGTTTC 1092
 ||||| ||||| ||||| ||||| |||||
1030 alHisGlnProSer.....GluGlnArgLeuGlnArgMet 1041
1093 AACACAGCGCTCAACGCGCGCGCGCGCATGTGCCGATTTGCTACTTA 1142
 ||||| ||||| ||||| ||||| |||||
1042 Gly.....AspThrGlyIylleHisAlaAspleuGlnIylsThrVa 1055
1143 CGAGCGC.....GTGATGCCCT 1159
 ||||| ||||| ||||| ||||| |||||
1055 lGlnuysTrpProAlaLeuAsnleuPheSerValAspHisValIylsAsnI 1072
1160 TGGATATCTCCGCCACCTGCTTTGCGCGATTAATGTCGCGCGATACC 1209
 ||||| ||||| ||||| ||||| |||||
1072 lGlnuAsnleuHisGlySerArgLeuAsnAspArgIleAlaGlyAspAsp 1088
1210 GACAGCGCGCGCATTTGGGT 1230
 ||||| ||||| ||||| ||||| |||||
1089 GlnAspAsnGlnleuTrpGly 1095
seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAU48589

seq_documentation_block:
ID AAU48589 standard; Protein; 372 AA.
AAU48589;
27-FEB-2002 (first entry)
Propionibacterium acnes immunogenic protein #9485.
SAPHO syndrome; synovitis; acne; pustulosis; hypertosis; osteomyelitis;
uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;
inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;
dermatological; osteopathic; neuroprotectant.
Propionibacterium acnes.
WO200181561-A2.
01-NOV-2001.
20-APR-2001; 2001WO-US12665.
21-APR-2000; 2000US-199047P.
02-JUN-2000; 2000US-208841P.
07-JUL-2000; 2000US-216747P.
(CORI-) CORIXA CORP.
Skelly YAM, Persing DH, Mitcham JL, Wang SS, Bhatia A;
L'maisonneuve J, Zhang Y, Jen S, Carter D;
N-PSDB: AAS59543.
WPI: 2001-616774/71.
Propionibacterium acnes polypeptides and nucleic acids useful for
vaccinating against and diagnosing infections, especially useful for
treating acne vulgaris -
Example 1; SEQ ID No 9784; 1069pp; English.
Sequences AAU39105-AAU68017 represent Propionibacterium acnes immunogenic
polypeptides. The proteins and their associated DNA sequences are used in
the treatment, prevention and diagnosis of medical conditions caused by
P. acnes. The disorders include SAPHO syndrome (synovitis, acne,
pustulosis, hypertosis and osteomyelitis), uveitis and endophthalmitis.
P. acnes is also involved in infections of bone, joints and the central
nervous system, however it is particularly involved in the inflammatory
lesions associated with acne vulgaris. A method for detecting the
presence or absence of P. acnes in a patient comprises contacting a
sample with a binding agent that binds to the proteins of the invention
and determining the amount of bound protein in the sample. The
polypeptides may be used as antigens in the production of antibodies
specific for P. acnes proteins. These antibodies can be used to
downregulate expression and activity of P. acnes polypeptides and
therefore treat P. acnes infections. The antibodies may also be used as
diagnostic agents for determining P. acnes presence, for example, by
enzyme linked immunosorbent assay (ELISA).
Note: The sequence data for this patent did not form part of the printed
specification, but was obtained in electronic format directly from WIPO
at ftp.wipo.int/pub/published_pct_sequences.
Sequence 372 AA;
XX

```

alignment_scores:      Length: 364
                        Ratio: 0.691
                        Gaps: 18
Percent Similarity:    43.132
                        Percent Identity: 22.253

alignment block:
US-09-303-518D-125/rev x AAU48589  ..

Align seg 1/1 to: AAU48589 from: 1 to: 372

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1280 GGCACACCAAGACGTCGACCAAGGAGGCTTCCTGCTCAATTCACGA 1231
    ||| : : : : : : : : : : ||| ||| : : : : :
21 GlyThrSerSerThrArgSerThrArgInsThrArgSerGlySerPro 37

1230 A.....C 1229
    :
37 TrpProSerSerValThrThrProValAlaArgThrCysLeuThrP 54

1228 CCAATGCTCGCGCGGCTG.....GTATGCCGACGATTTAA 1191
    || : : : : : ||| : : : : : ||| : : : : :
54 TrpLeuGlySerGlyThrSerProSerProProSerProArgLeuSer 70

1190 TCGCGCAAAAGAGGGTGGGAGATATCCAAAGGACATCAGCGCTGCTA 1141
    ||||| : : : : : : : : : : : : : : : :
71 SerArgLeuAlaThrThrProValLeuThrThrAlaLeuAlaSerThrVa 87

1140 AGTACCAATCGGCGACCATATGCGCGGCTGCCCGCTTGACGGCTGTGTA 1091
    : : : : : || : : : : : ||| : : : : :
87 LProthrValAlaGlyThrSerSerSerLysProGlyProSerCys... 102

1090 ACTTGAAAGATTGTGTTTTCAGAAATAGGCGAGGGTGTATGCGGTATG 1041
    : : : : : |||||
103 .....TrpProPro..... 105

1040 GAGTATTTTTCGGGCTCGCGGCGCAACCCAGCGCAACGCTTTTGCTGC 991
    : : : : : : : : : : ||| : : : : : : :
106 .....AlaAlaProSerSerLeuThrProThrThrSerAlaIleSe 119

990 GCCTTCTTTCG..... 981
    |||||
119 rThrSerSerAlaProProSerProLeuArgCysThrGlySerArgCysA 136

980 .....ATAACGAAATCTGATTTGTGTAGCGTCCCAATAATATCGTGC 939

136 TGCystrPheuThrArgLeuLeuAlaGlyThrThrAla.....ProAla 150
    : : : : : ||| ||| : : : : : : :
938 GCGCGCTTGATGTATGCGCGCGCTCAATATCGCAACCGGAATCAGCGGTT 889

151 SerProGlySalAlaVal...ProSerValThrSerProThrLeuLeuGly.. 165
    : : : : : : : : : : ||| : : : : : : :
888 GTCTGTGTCAACAACATTCGCC.....GCATATATTTGGCATA 851
    : : : : : : : ||| : : : : : : :
166 GlyGlyLysnGlySancysProProAlaProSerSerValValThrAsp 182

850 CTTTCGCACCCCAAAACGGTACGCAAGAGCGCGGTGTGTTGACTTGAAGA 801
    : : : : : ||| ||| ||| : : : : :
182 LysSerSerPro.....ArgProArgSerGlySerSerAlaProAsp 195

800 CCACCTTAGCGCAATCAGCGCGTCTGCTTACAGACGCGCTTCGCAACAA 751
    ||||| : : : : : : : ||| |||
196 ProProLysasnThrArgInsThrArgAlaArgProPro..... 208

750 ACGGCCAATGCTAATTACATCTGTGATTAATTGATGCTCCACAGGTTTTAT 701
    ||| ||| : : : |||
209 SerProProValSerSerSer..... 215

700 TCGGCGCAGCCGCGCTCGATGTAATGATGTGCGTCCACATCAACCGGCA 651
    : : : : : ||||| : : : : :
216 .....LeuLysProSer 219

650 GGATCGCGGCGCGCAATTCATGTGTTTCGATGTGCGACACATTTTCAGA 601
    : : ||| : : : |||||
220 ...TrpSerProAlaArgTrpThrValSer..... 228

600 CGGACAGCTTCGCGCACCTGCTTACAAACATGGAATTTTGCTTCGGTCA 551
    ||| ||| : : : : ||| ||| : : : : :
229 ....ProSerLeuProSerAlaLysProThrSerThrArgArgAlaThrS 244

550 AACGGCTCAATACCAACAGCGCGGTGTGAAATCTCGCGCGGCTCTTTGG 501
    : : ||| : : : : ||| : : : ||||| : : :
244 eraGlyTrpArgSerSerAlaProGlySerGlyThrSerAlaThrValGly 260

```

seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AAL1998.DAT: AAM80699

seq_documentation_block:
ID AAM80699 standard; Protein; 835 AA.
XX
AC AAM80699;
XX
DT 24-DEC-1998 (first entry)
XX
DE S. pneumoniae cation transporting ATPase.
XX
KW Streptococcus pneumoniae protein; recombinant; gene expression; DNA chip;
KM virulence; antibody; infection; detection; treatment; hypothetical;
cell wall biosynthetic, external target; minimal gene set protein.
XX
OS Streptococcus pneumoniae.
XX
PN MO9826072-AI.
XX
PD 18-JUN-1998.
XX
PE 09-DEC-1997; 97MO-US22578.
XX
PR 13-DEC-1996; 96US-0036281.
XX
ELIL) LILLY & CO ELI.
PA
PI Baltz RH, Burgett SG, Dehoff BS, Hoskins JA, Jaskunas SR;
PI Mills BJ, Norris FH, Peery RB, Rockey PK, Rosteck PR;
PI Skatard PL, Smith MC, Solenberg PJ, Treadway PJ;
PI Young Bellido ML;
XX
DR WPI: 1998-348529/30.
DR N-PSTD: AAV65261.
XX
PT Streptococcus pneumoniae nucleic acid sequences - used in DNA chips
PT for evaluating gene expression, and identification of virulence
PT genes
XX
Claim 3; Pages 288-291; 333pp; English.

This sequence represents a Streptococcus pneumoniae cation transporting ATPase. The invention provides DNA sequences (AAV65201 to AAV65304) from the Streptococcus pneumoniae genome and corresponding protein sequences (AAM80605 to AAM80728). The protein sequences are classified as hypothetical, cell wall biosynthetic, external target, or minimal gene set proteins. A recombinant host containing a vector comprising any of the above nucleic acids can be used for the recombinant expression of the proteins. The invention also provides a DNA chip having arrayed on it at least 15 base pair fragment of any one or more of these DNA sequences. The DNA chip can be used methods for evaluating gene expression in S. pneumoniae and for identifying virulence genes in S. pneumoniae. Antibodies that selectively bind to the above proteins or peptide fragments can be used to treat S. pneumoniae infection. The antibodies can also be used to detect S. pneumoniae cells.

Sequence 835 AA:

alignment_scores:	Quality:	108.00	Length:	356
Ratio:	0.617	Gaps:	15	
Percent Similarity:	49.157	Percent Identity:	23.315	
alignment_block:				
US-09-303-518D-125 x AAW80699 ..				
Align seg 1/1 to: AAW80699 from: 1 to: 835				
28	CTGGCCATCGCGGCGACACCGGACGCAAGCGCTTACGACGGCCCGGCAT	77		
224	LeuAlaValAlaAlaAlaIleProGluGlyLeu.....ProAlaI	236		
78	TACGCAATCGCGCTTG...CTTGGCGAAGATATATGCGCGATCGCCCT	124		
236	evalThrlleValleuSerLeuGlyThlGlnValleuAlaLysArgHis	253		
125	CGATG.....AAGTCAGAGAGCGCATGCCCGTCAAAAAGCCCAAGTG	168		
253	erlleValArgLysLeuProAlaValGluThlThleuGlySerThrluile	269		
169	CTGTGTGAAGCAAAAGAAAT.....CC	191		
270	lleAlaSerArgLysThrGlyThlLeuThlMetAsnLysMetThlValGl	286		
192	GGGCGGTGTGTTCCTGCGCGGCTTACGCAAAATCGCGCGATTCAC	241		
286	uLysValIlePheArgLysAlaValAlaLeuHisArgSerAlaAspArgIleGlu	303		
242	GTGCG...GAAAAGCGGCTACTGATGATGCTGATGCGCGCTGACAGC	288		
303	eugLysLeuGluMetArgProLeuArgSerValValLeuAlaAsnSprThr	319		
289	AACACAGCAATTCGATTTGAAGCGCTACGCGCTGAAAGCGCTGCAAACTT	338		
320	LysIleAspValGlu.....GlyAsnLe	327		
339	AAGCGGCGAAGAGTGGCGCGGACCTGATCCAAATCGGTTGTGTGACAG	388		
327	uIleGlyAspProThrGlnThlAlaPheIleGlnThlAlaLeuAspArgLysG	344		
389	CGGTGCGGACCGGCGCTTG...ACGAAATTCGCGCGCTGATGCGCGAG	435		
344	LYTYAspValLysGlyPheLeuGluLysGlyProArgAlaAlaGluLeu	360		
436	CGGTGCGGCATTTGTCATATGCGATGCGACCC...AATCGGTGCGTCC	482		
361	ProPheAspSerArgLysLeuMetSerThrValHisProLeuProAs	377		
483	CGACCTTAGCGCATATTCAAAGAGCGCGCGAGATTCCAAAGCGGCGC	532		
377	PserArgPheLeuValAlaValLysGlyAlaProAsp.....	389		
533	TGTTGTATAGACCGGTTGACCGGCAAGAAATCCATGTTGTATAGCA	582		
390GlnLeuLeuLysArgCysLeuLeuArgAspLysAla	401		
583	GCTGCGCGACAGCGCGCTGTGAAATGCTGCCAAC...ATCGAAACACA	629		
402	GlyAspIleValIleArgProIleLeuLysValIleThrAsnLeuIleHisThrAs	418		
630	TGAATTTGGCGCGCGCATCTGCGC.....GGTTGAGTGGGACGACACA	673		
418	msnSerGluMetAlaHisGlnAlaLeuArgValLeuAlaGlyAlaLysArg	434		
674	TTTCATTATCGAGCGCGTGGCGCGCAATTAACCGCTGTGACCATCAAT	723		
435	..LysIleIleAspSerIleProGluAsnLeuThrSerGluGluLeuGlu	450		
724	TATCAAGATGTATTCATTCATGGCGGCTTGTGTGCAACAGCGCGTGTGAA	773		

Align seg 1/1 to: ABG23389 from: 1 to: 1194

```

1313 TTGCGCAACAGCGGGCGCTATTGCTATTGATTTGCCGGGAGACAGACACTGCA 1264
    ||| :|||:||||| :|||:|||||
584 LeuSerGlnSerGlyProProGlyLeuLeuPro..... 594
1263 CAAGAGGAGGCTCTTCTGCTCAATTCAGACACCAATGCC...TCGG 1217
    ||| ||| :|||:||||| :|||:|||||
595 .....SerProSerPheAspSerIysProProThrThrLeuLeuG 608
    :||| :|||:|||||:
608 ILeuIleProAlaProSerMet..... 615
1166 AATATCAAGGGGATCATCGCGCTGTAAGTACCAATTCGACCACTAGCGCG 1117
    ||||| :|||:||||| :|||:|||||
616 .....ValPro..AlaThrAspThrIly 622
1116 GTGCGCGCGCTTGAGCGCTGTTGACTTAAGAGATTGTTTTCAGGA 1067
    :|||:||||| :|||:||||| :|||:|||||
622 SalAProProThrLeuGlnAlaGlnThrAlaThrIlyProGlnAlaThrS 639
1066 AATGCGCCGAGGGTTGTACGCGTGATGAGATTTGTCGGGCGGCGCA 1017
    :||| :||| :|||:|||||:|||||
639 erAlaProSerProAlaProIylsGlnSerPheLeuPheGlyThrGlnAsn 655
1016 ACCAGCGCGAAGAGCTCTTGTGCGGCTCTTTCGATTAACGGAATCTG 967
    |||||:||||| :|||:||||| :|||:|||||
656 ThrSerProSerSer.....ProAlaAla..... 663
966 ATTGTGTAGGAGTCCCAATATATGTCGGCGCTGTGTATGCGCGCT 917
    ||| ||| ||||| :|||:||||| :|||:|||||
664 .....ProAlaAlaSerSerAlaProIlePheIlyProI 676
916 TCAATACCGAAGCGGAATACGCGGTTGTGTGTCAACCAATTCGCC 867
    :||| ||| :|||:|||||:
676 IePheThrAlaPro..... 680
866 GCATATATTGCGATCTTTCGACCCAAACGCTACGCAAGAGCGCGG 817
    |||||:||||| :|||:||||| :|||:|||||
681 .....ProIylsSer.....GlnIylsGlnI 687
816 TTTGTGACTTGAAGAACCACTAGCGCAATCAACGCGGTGTAGAC 767
    ||| :||| :|||:|||||:|||||
687 yProThr.....ProProGlyProSerValThrAlaThrAla..... 699
766 GCGCTGTGCAACAAACGCGCAATGATTAATTCATCTTGATTAATGATG 717
    ||| :|||:|||||:|||||
700 ..ProSerSerSerLeuProThr..... 708
716 GTCCACAGCGTTTATTCGCGCGGCGGCTGATGAATGAATGCGCT 667
    ||| |||||:|||||:
709 .....ThrSerThrThrAlaProThr..... 715
666 GCCACTCAAAACG.....GCAGGATGCGGCGCGCAATTCATGTTT 623
    :|||:||||| :|||:||||| :|||:|||||
716 ....PheGlnProValPheSerSerMetGlyProProAlaSer...ValP 730
622 CGATGTGGCAGCATTTTCAGACGCGAGCTGTGCGCGACCTCCCTTACAA 573
    :||| :||| :|||:|||||:|||||
730 roLeuProAlaProPhePheIylsGlnThrThrProAlaThrAlaPro 746
572 ACATGATTTTGCCTGCGTCAACGCGCTCATACCAACAGCGCGCTT 523
    ||| :||| :|||:|||||:
747 Thr..... 747
522 GAAATCTGCGCGCTCTTTGATATATACCGTAGGTCGCGACGACGCG 473
    :|||:||||| :|||:||||| :|||:|||||
748 .....ThrThrAlaProLeuPheThrGlyLeuAlaSerAlaThrSera 762
472 GATGCGTGCATGCATTCAGAGAGATGGCAACGCGCTCGCATCG... 426
    :|||:||||| :|||:||||| :|||:|||||

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762 IValAlaProIleThrSerAlaSerProSerThrAspSeraIaSerIys 778
425 ...ACGGCAGGAATTTGCTGAACGAGCGGTGTCGCGACCATCCACAA 379
    :|||:||||| :|||:||||| :|||:|||||
779 ProAlaPheGlyPheGlyIleAsnSer...ValSerSerSerValSera 794
378 ACCGATTCGATCAGGTTGGCGGCACCTTCTTGCGCGCTTAAGTTGCCA 329
    :|||:||||| :|||:||||| :|||:|||||
794 rThrThrThrSerThrAlaThrAlaIaSerGlnProPheLeuPheGlyA 811
328 GC.....GCTTCAGGTGCGTACGCTTCAAACTCGATTGCGTGTG... 288
    :|||:||||| :|||:||||| :|||:|||||
811 IapProGlnAlaSerAlaIaSerPheThrProAlaMetGlySerIlePhe 827
287 .....CCTTCACGCGCAATCACGACTGACTGAAGTACGCGCTT 250
    :||| :||| :|||:|||||:|||||
828 GlnPheGlyIylsProProAlaLeuProThrThrThrValThrThrPh 844
249 TTGCGCAGGTGAATGCGGCGATTTTGCTGAAGCGCGCGCATTAACA 200
    ||||| :|||:||||| :|||:|||||
844 eSerGlnSer...LeuAlaThrAlaValProThrAlaThrSerSerSera 860
199 CCAAG.....CCCGATTCCTTTGTCTTCAACAGCACTTGCGCTTTT 156
    :|||:||||| :|||:||||| :|||:|||||
860 IAlaAspPheSerGlyPheGlySerThrLeuAlaThrSerIaProAla 876
155 TTGACGCGATCCGCTTCTGACTTCATCGAGGCGCATACGCGCAT 106
    :|||:||||| :|||:||||| :|||:|||||
877 ThrSerSerGlnProThrLeuThrPheSerAsnThrSerThrProThrPh 893
105 TTCTTCGCGCAACAGCAACGCACTTCGTAATGCGCGGCGCGCTGAACG 56
    :|||:||||| :|||:||||| :|||:|||||
893 eAsnIleProPheGlySerSerAlaIylsSerProLeuProSerIylsProG 910
55 CTTCGCTCCGCTGCGCGCGATGCGC 30
    :|||:||||| :|||:||||| :|||:|||||
910 IylAlaAsnProGlnProAlaPheGly 918

seq_name: /SIDSI/gcgdata/geneseq/geneseq-emb1/AA1197.DAT:AAW13504
seq_documentation_block:
ID AAW13504 standard; protein; 1644 AA.
XX
AC AAW13504;
XX
DT 30-JAN-1998 (first entry)
XX
DE B. bronchiseptica adenylcyclase-haemolysin mutant delta-Cla.
XX
KW Mutant; Bordetella pertussis; adenylcyclase; haemolysin; wild type;
KW deletion; induction; antibody; Bordetella parapertussis; vaccine; human;
KW Bordetella bronchiseptica; veterinary.
XX
OS Bordetella bronchiseptica.
XX
FH Synthetic.
XX
FH Key Location/Qualifiers
FT Region 826..827
FT /note= "deletion of amino acids 827-887 from the wild
FT type sequence between these positions"
FT Modified-site 919..924
FT /note= "Arg at position 922 is modified by addition of
FT a fatty acid"
XX
XX FR2736064-AL.
XX
XX PD 03-JAN-1997.
XX
XX PD 30-JUN-1995; 95FR-0007945.
XX
XX PR 30-JUN-1995; 95FR-0007945.
XX
XX PA (INSP ) INST PASTEUR.

```



```
XX AC AAY04954;
XX DT 06-JUL-1999 (first entry)
XX DE Mycobacterium species protein sequence 41T#2.
XX KW Secreted protein; Mycobacterium; primer; PCR; amplification; probe;
XX KM hybridisation; detection; vaccine; immunisation; infection.
XX OS Mycobacterium sp.
XX PN WC0909186-A2.
XX PD 25-FEB-1999.
XX FE 14-AUG-1998; 98WO-FR01813.
XX PR 11-SEP-1997; 97FR-0011325.
XX PR 14-AUG-1997; 97FR-0010404.
XX PA (INSP ) INST PASTEUR.
XX PI Glacquel B, Lim EM, Pelicic V, Portnoi D, Goguet de la Salmoniere Y,
XX PI Guigueno A;
XX DR WPI: 1999-181045/15.
XX DR N-PSDB: AAX34206.
XX PT Mycobacterial DNA vectors containing reporter constructs - for
XX PT identifying coding or promoter sequences involved in
XX PT infection-associated protein expression
XX PS Claim 32; Fig 41T; 309pp; French.
XX CC Sequences AAY04742-Y05000 and AAY07201-Y07204 represent secreted
XX CC proteins from various Mycobacterium species microorganisms. The
XX CC encoding nucleotide sequences can be used as primers and probes for
XX CC methods for detecting and identifying mycobacteria, especially belonging
XX CC to the M. tuberculosis complex. The encoded proteins can be used in
XX CC vaccines for immunisation against a bacterial or viral infection.
XX SQ Sequence 572 AA;

alignment_scores:
      Quality: 106.00      Length: 509
      Ratio: 0.586      Gaps: 33
      Percent Similarity: 35.560      Percent Identity: 20.825

alignment_block:
US-09-303-518D-125 x AAY04954 ..

Align seg 1/1 to: AAY04954 from: 1 to: 572

26 ACCTGCCATCGCGGACGACGCGGATTCAGACGCGCGCC 75
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   :|||
113 SerCysProArgSerCysAlaGlySerGlnArg**ProArgLeuArgPr 129
   :|||
   :|||
76 ATTACCGAAGTCCGCTTGGTGGCGAAGATATGCGGTA...TGGCGCC 122
   :|||
   :|||
129 oProProLeuAlaArgItyrCysGlyArgSerThrProThrProSerGlyP 146
   :|||
   :|||
123 CTGATGAAGAATCAAGAGGCGATCGCTCAAAAAG..... 160
   :|||
   :|||
146 roArgGlyGlyAlaAlaSerSerThrProThrProValAlaArg 162
   :|||
   :|||
161 .....GCCAAGTGTCT.....TT 174
   :|||
163 ArgCysArgGluLeuSerSerArgCysGlyProProGluProSe 179
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175 GAAGAGAAAAGAAATCCGGCGTGGTTTACTGCGCGGCTTCAGGCA 224
   :|||
   :|||
```

```
179 rThrsAlaArgThrArgGlyTrp.....ProVal 190
225 AATGCCCGCATTCACCGTGGCGAAGACCGTACTTCACTCAGTCGTGA 274
   :|||
   :|||
190 rSerProProProValThrArgArgSerSerArgTrpLysArg..... 204
   :|||
   :|||
275 TTGGCGTTGAAGCAGCAGCAATGAGTTTACAGCTACGACCTGAA 324
   :|||
   :|||
205 ...ProProAsnSerLeuThrCysSerProMetArgAla..... 216
   :|||
   :|||
325 GCGGTGGCAAACTTAAAGCGCGAAGAGTGGCGCGACACCTGATCCATC 374
   :|||
   :|||
217 ArgTrp...ThrProAlaSpGlyAlaCysTrp..... 226
   :|||
   :|||
375 CGGTTGTGATGCTGCGTGGCGACCGCTCCGTTCAAGAAAATTCTGCG 424
   :|||
   :|||
227 ...PheCysTrpThrArgCysAlaProProSerAlaGlyArgHisLeuPro 242
   :|||
   :|||
425 TCGATGCCGAGCCGTTGCCATCTGTCATGATGATGACCAATCCG 474
   :|||
   :|||
243 GlyArg...SerThrAsnProArgArgAla...ArgCysArgProThrAr 257
   :|||
   :|||
475 CTGCTGCCGACCCCTACGGTCAATTATCAAAAGACCGCGAGATTTCAA 524
   :|||
   :|||
257 g.....LeuProAsnAlaProProArgAsnSerA 267
   :|||
   :|||
525 ACGCGGCG...TGTGTATGTAGCGCGTTGACCGACCAAAATTCATG 571
   :|||
   :|||
267 rg**CysIleCysTrp..... 272
   :|||
   :|||
572 TTTGTAAAGCAGCTGGCGACGACGTCGCGCTGAAATGTCGCAACATC 621
   :|||
   :|||
273 .....ArgTyrValMetLeu.GlnArgA 280
   :|||
   :|||
622 GAACACATGAATTGGCGCGCGCCGATCTGCGGTTTGAG..... 662
   :|||
   :|||
280 rg.....ThrsCysGlyIleAspSerArgAsn 289
   :|||
   :|||
663 TGGACGACATTCATTTCATGACGCGGTGGCGGCAATAAACCGTGT 712
   :|||
   :|||
290 Trp..... 290
   :|||
   :|||
713 GGACCATCATATATCAAGATGTAATTACCATGCGCGTTGTTCACAA 762
   :|||
   :|||
291 .....ValSerArg.....TrpProSerProLeuAla 300
   :|||
   :|||
763 GCGCGTGTGAACACGACGCGCGTGAATGCCCT..... 794
   :|||
   :|||
300 rgProThrAlaThrProItyrThrSerThrProThrThrProValProPro 316
   :|||
   :|||
795 .....AGGTGGTTCATCAATCAACAAACCGCGCTCTTGC 829
   :|||
   :|||
317 TrpLysProAspTrpArgTrp.GlyGluLeuAlaGlySer**SerArgA 333
   :|||
   :|||
830 GTACCGTTTGGGTGCGAAGATATGCAAAATTAATCTGCGGGAATGGTT 879
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333 rgSerValProGly..... 337
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   :|||
880 GACACAGACAAACCGCGTGAATTCGCGTTGCGTATTTGAAGCGCGGATTC 929
   :|||
   :|||
338 ProAlaAspCysArgProValAlaGly..... 346
   :|||
   :|||
930 ACAAGCGCGACGATTAATTGGGACCTACCAACATCAGATTTCGCTTA 979
   :|||
   :|||
347 ArgGlyAlaAlaProCysTrpArgSerSerThrAlaThrValPro... 361
   :|||
   :|||
980 TCGAAGAGAGCGCGACGCAAGAGCTGTTCGCTGGGTGGCGCGCAC 1027
   :|||
   :|||
362 .....ProSerCysSerProGlyArgAlaProAlaLacy 372
   :|||
   :|||
1028 .....CGGCAATATCCATCAGCGGTACAAACCTCGCGCATTTCTT 1070
   :|||
   :|||
372 scysAspArgValAlaThrPro**HisArgProProIleSerVal.... 387
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```

1071 GAAAGCAACTCTTCAAGTTCAACACAGCGGTCACGGGCGACCGCG 1120
      :::::  :::::  :::::  :::::  :::::  :::::
388 .....ProthSerTrpCysGlyProtrp**ThrProAlaProArgThr 402
1121 CCATGTCGCCGATTGTTACTTACGAGCGCGATGCCCT..... 1159
      |||||  |||||  |||||  |||||  |||||  |||||
403 ***TrpCys.....CysProMetAlaMetTr 411
1159 ..... 1159
411 pProProlAsnTrpTrpProGlyValProArgArgSerAlaGlyAlas 428
1160 .....TGATATCTCTGC...CCACCCGCTTTTCCGGATTATCTCTGC 1201
      |||  |||  |||  |||  |||  |||
428 ertHrTrpTrpCysArgProAspArgTrpCys..... 439
1202 GCGATACGACAGCGCGCATTTGGCTTGTGAATTTGACGAGAA 1251
      |||  |||  |||  |||  |||  |||
440 .....ArgGlyTrpProArgTrpPro..... 446
1252 GACCTGCTTTTGTGACCTTGTCTGCTGCCGCGCAATACG..... 1291
      |||  ::::  |||||  :::::  |||
447 .....CysMetThrArgProAlaArgProSerThrThrAlaThr 460
1292 .....AATAGCGCCGCTGTGGCAAG 1315
460 lATrPProValProProValLeuProGlyThrAspArgCysAlaLeu... 475
1316 TGCTGAAACCATTTGAGAAG 1336
      |||  |||||
476 .....ProProlArgArg 480

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seq_name: /SIDS1/gcgcdata/geneseq/geneseq-emb1/AA2001.DAT: AAB59814
seq_documentation_block:
ID AAB59814 standard; Protein: 1022 AA.

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```

XX AC AAB59814;
XX DT 04-APR-2001 (first entry)
XX DE Tugd protein #5.
XX KM Toluene degradation; enzyme; waste degradation; Tugd.
XX OS Thauera aromatica.
XX OS Xanthomonas maltophilia.
XX OS Geobacter metallireducens.
XX OS Azarcus tolulyticus.
XX PN W020072650-A2.
XX PD 07-DEC-2000.
XX PF 24-MAY-2000; 2000WO-US14298.
XX PR 01-JUN-1999; 99US-0323872.
XX PA (UYOH-) UNIV OHIO.
XX PI Coschigano PW.
XX DR WPI: 2001-041080/05.
XX DR N-PSDB: AAF23625, AAF23627.
XX PT Composition comprising toluene degrading enzyme useful for biological
XX PT treatment of organic compounds, especially for degrading toluene or its
XX PT analogs
XX PS Disclosure: Fig 5; 122pp; English.
XX CC The present invention relates to toluene degrading enzyme genes and

```

CC proteins tuth (see AAF23629 and AAB59831), tuti (AAF23630 and AAB59832), CC tuf (AAF23631 and AAB59833) and tulg (AAF23632 and AAB59834). The CC toluene degrading enzymes are homologues of private formate lyase. The CC toluene degrading enzymes are useful for biological treatment of organic compounds and in particular for the degradation of toluene and its CC analogs contained in liquid or solid waste source. The present sequence is a protein sequence for toluene degrading enzyme, Tugd.

Sequence 1022 AA;

alignment_scores:
 Quality: 106.00 Length: 498
 Ratio: 0.535 Gaps: 27
 Percent Similarity: 39.759 Percent Identity: 24.096

alignment_block:
 US-09-303-518D-125 x AAB59814 ..

Align seg 1/1 to: AAB59814 from: 1 to: 1022

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18 AGGCTTAACCTGCCATCGCGGCGAGACCGGAGCAAGCGTTTA..... 62
      ||:::  |||  ||:::  ||:::  ||:::  ||:::
19 ArgThrValProHisGlyArgAlaGlyLeuProAlaGlyArgLeuAl 35
63 .....CGACGCCCGGCCATTACGAAAGTCCGTTGCTGGCGAA 102
      |||||  |||
35 aAlaAlaGlyArgArgGly.....GlyAsp 45
103 GAATATGCGCGGTATGCGCCCTCGATGAAGTCAGGAAGCGATCCGT 152
      ::::  ::::  |||||  ::::  ::::  |||
45 lIleLeuGlnAlaAlaProAla...GlnIleValSerAlaLeuPheArg 60
153 CAAAAAGGCCAAGTGTGTTGAAGCAAAAGAAATCCGGCGGTGTGT 202
      :::  |||||
61 SerGlyArgProArgProHisValSerGlyGlnGlnHisGlyAlaVal 77
203 TTACTGCGCGGCTTCAGCAAAATCGC.....CGGATTCACCGT 243
      |  :::::  ||:::  |||
77 lLeuArgPheArgLeuGlnGlnHisArgAlaAlaLeuArgAsnArgPro 94
244 GCGGAAAAAGCGCTACTTCAAGTCAGTCGTGATTCGCGTTGAGCAACA 293
      ||:::  |||
94 lArgArgAla..... 97
294 CGAATGAGATTGAAGCTACGACCTGAAGCGGTGCAAACTTAAGCG 343
      |||||
98 .....AlaGlyAlaHisGlnAl 103
344 GCGAAGATGCGCGCGCAACCTGATCCAAATCCGTTTGTGACTCGCTG 393
      |||||  ::::  |||  |||||
103 aArgArgLysAlaGlyArgArgSerProGlyArgHisValAspLeu... 118
394 CGCACCGCGCTCCGTTACACAAATTCCTCGGTGATGCCGACCGTTCCG 443
      |||  :::::  ||:::  |||
119 .....ProLeuGlnArgHisGlnGlySerArgGlnHisArgGlnAspArg 133
444 CATCTGCTCAATGCGATGAGACCAATCCGCTGC.....TG 481
      :::::  |||  |||||  ::::
134 GlnLeu...GlnGlyAspGlyHisArgLeuGlnGlyAspGlnLeuG 149
482 CGGACCTACGCTATTATCAAGAGCGCGCGAGATTTCAAACGCGCG 531
      |||  |||  :::::  |||||
149 yAlaProAlaGlyProAlaValGlnAspArgArgGly..... 161
532 CTGTTGGTATGACCGCTTACCGAAGCGCAAAATTCATGTTGTAAAGC 581
      :::  |||||  :::::  |||
162 .....LysLeuArgAspArgSerGlnAlaPro..... 170
582 AGTGGCGCAGACGTCCGCTGAATAATGTCACACATCGAACAACATG 631
170 ..... 170

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632 AATTCGGCGCCGCGATCCTGCGGTTTGTAGTGGACGACATTCATTC 681
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171 ....GlyArgAlaAlaArgAsnArgHisLeuSerAlaHisSer...A 185
682 ATGACCGCGGTGGCGGCAATAAACCGTGGACCATTAATTCACAGA 731
    :||||| :||| :||||| :||||| :||||| :||||| :|||||
185 rgrArgAlaLeuGlnGlyProGlnGlyArgAspAlaGlyIle... 199
732 TGTAAATACCATTGCGGTTGTTTGCAC... 761
    :||| :||| :||| :||| :||| :||| :||| :|||
200 ....LeuTyrLeuProAspLeuSerArgAspArgAlaLeuArgGln 214
762 ....AGCGG...TCGAACACCGACCGCGGTATGCCCTAGCGTCT 804
    ||||| ||||| ||||| ||||| ||||| ||||| |||||
214 gLeuArgProGlnGlyArgHisProAlaValAlaValLeuGlnGlyLeu 231
805 CAAGTCACAAACCGCGCTTGGCGTACCGCTTGGGCG... 845
    :||| :||| :||| :||| :||| :||| :||| :|||
231 rArgArgGlnGlnIleProAlaAspGlnProHisGlyGlyGlyThr 247
846 .....GAAAGTATCGCAATTACTCGGCGCAATTG 877
    ||||| ||||| ||||| ||||| ||||| ||||| |||||
248 ArgArgAspGlyThrPheGlnAspPheArgAlaTrpArgGlnValAl 264
878 TTGACACAGACAAACCGCGGATTCGCGGTGCGTATGCA... 917
    :||| :||| :||| :||| :||| :||| :||| :|||
264 a.....ArgLeuProArgAsnLeuPro...GlyValGlnArgSerVal 278
918 .....CGGCGGATTCACAAAGCGCGACGATTTATGGGACGCTA 959
    ||||| ||||| ||||| ||||| ||||| ||||| |||||
278 IsProHisArgArgHisGlnArgGlnGlyArg....GlyArgLeu 292
960 CCACATATGATTCCTTATCGAAGACGCGACGCAAGAGCTGTTCG 1009
    :||| :||| :||| :||| :||| :||| :||| :|||
293 GlnArgHisAspArgArgHisProArgGlySerGln... 304
1010 COTGGTGGCGCGCGACCGCAATCTCCACAC... 1046
    :||| :||| :||| :||| :||| :||| :||| :|||
305 .....AlaAspProHisGlyArgAlaLeuHisArgLeuProLeuPhe 319
1047 .....GCGTACAAACCTCGCGCATTCCTGTGAA 1073
    ||| ||| ||| ||| ||| ||| ||| |||
319 GlnGlnLProArgGlnAspAlaAlaLeuGlyPheArgValHisProArg 335
1074 AAACAACTCTCAAGTTCACACACGCGCTCACGCGCGACGCGCA 1123
    :||| :||| :||| :||| :||| :||| :||| :|||
336 ArgThrArgLeuSerValAsp.....GlnAlaArgArgAspArgHis 349
1124 TGGTGGCGATGTACTTACGAGCGCGTATGCCCTTGATATCTGCC 1173
    ||||| ||||| ||||| ||||| ||||| ||||| |||||
349 sGlyAlaAspGlnGlyIleArg.....GlnValG 359
1174 ACCCTGCTTTGCGGATTAATCTGCGGATACCGACGCGCGAGCG 1223
    :||| :||| :||| :||| :||| :||| :||| :|||
359 LPro.....GlnArgGlnArgArgHisAlaArgHisProArg 366
1224 ATTGGTGTCTTGAATTGCA.....CGAAGAAGACCTGCTT 1261
    ||||| ||||| ||||| ||||| ||||| ||||| |||||
370 SerProGlnLeuGlnArgAlaValHisValAlaArgHisProArg 386
1262 TGTGCA.....CTTCGCTGCTGCGG 1281
    ||| ||| ||| ||| ||| ||| ||| |||
386 rProGlnAspAlaLysAsnProPheGlyArgTrpArgLeuAsnLeuPro 403
1282 GCGCAATACGAATACGCGCGCTGTGCGCAAGTCTGGAAC 1325
    ||| :||| :||| :||| :||| :||| :||| :|||
403 Ly.....GlnAlaAlaGlyAsn 408
seq_name: /SID1/gcgdata/geneseq/geneseq_emb1/AA2001.DAT.AAB59824
seq_documentation_block:
ID AAB59824 standard; Protein: 1605 AA.
XX

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AC AAB59824:
XX
XX 04-APR-2001 (first entry)
XX
XX Protein #1 encoded by TdtD/E gene.
XX
XX DE
XX
XX KM Toluene degradation; enzyme: waste degradation; TdtD.
XX
XX OS Thauera aromatica.
XX OS Xanthomonas maltophilia.
XX OS Geobacter metallireducens.
XX OS Azotobacter toluyticus.
XX
XX PN W0200072650-A2.
XX
XX PD 07-DEC-2000.
XX
XX PF 24-MAY-2000; 2000WO-US14298.
XX
XX PR 01-JUN-1999; 99US-0323872.
XX
XX PA (UOH-) UNITV OHIO.
XX
XX PI Coschigano PW;
XX
XX DR WPI; 2001-041080/05.
XX
XX DR N-PSDB; AAF23627.
XX
XX PT Composition comprising toluene degrading enzyme useful for biological
XX treatment of organic compounds, especially for degrading toluene or its
XX analogs
XX
XX PS Disclosure: Fig 12; 122pp; English.
XX
XX CC The present invention relates to toluene degrading enzyme genes and
XX proteins tuth (see AAF23629 and AAB59831), tdtI (AAF23630 and AAB59832),
XX tufE (AAF23631 and AAB59833) and tufG (AAF23632 and AAB59834). The
XX CC toluene degrading enzymes are homologues of pyruvate formate lyase. The
XX CC toluene degrading enzymes are useful for biological treatment of organic
XX compounds and in particular for the degradation of toluene and its
XX CC analogs contained in liquid or solid waste source. The present sequence
XX is a protein sequence encoded by toluene degrading enzyme gene, TdtD/E.
XX
XX SQ Sequence 1605 AA;

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alignment_scores:

	Quality:	106.00	Length:	498
	Ratio:	0.535	Gaps:	27
	Percent Similarity:	39.759	Percent Identity:	24.096

alignment_block:

US-09-303-518D-125 x AAB59824 ..

Align seg 1/1 to: AAB59824 from: 1 to: 1605

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18 AGCTTAACCTGCGCCATCGGCGGACGACCGGACGACCCGTTA..... 62
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603 ArgThrValProHisGlyArgAlaGlyLeuProAlaGlnArgLeuAl 619
63 .....CGACGCGCGCGCATTAACCGAAGTGGCTTGGCGAA 102
    ||||| ||| ||| ||| ||| ||| ||| |||
619 AlaAlaGlyArgArgGlyGly.....GlyAsp 629
103 GAATATGCGCGGTATGCGCGCTCGATGAAGTCAAGAGGCGATGCCGT 152
    :||| :||| :||| :||| :||| :||| :||| :|||
629 InLeuGlnAlaAlaProAla...GlnGlnValSerAlaLeuPheArg 644
153 CAAAAAGGCCAAGTGGCTTTGAAGACAAAAAATCCGCGCGCTGT 202
    :||| :||| :||| :||| :||| :||| :||| :|||
645 SerGlyArgProArgProHisValSerGlyGlnGlnHisGlyAlaVal 661
203 TTACTGCGCGCGCTTACGCGCAAAATCGC.....CGGATTCACCGT 243

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661 1LeuArgPheArgLeuGlnGlnHisArgAlaAlaLeuArgAsnArgProG 678
      :::::::::::::::::::: |||::: |||
244 GCGCAAAAGCGCGTACTTCACTGAGTCGTGATTCGCGTTGAAGCACA 293
      |||::: |||
678 1ArgArgAla..... 681
294 CGAAATGAGTTTGAACGCTACGACCTGAAGCGCTGGCAAACTTAAGC 343
      |||::: |||
682 ..... 687
344 GCGAAGATGGCGCGCAACCTGATCCAACTCGTTGTGAGCGCGCTG 393
      |||::: |||
687 ArgArgLysAlaTyrArgArgSerProGlyArgHisValAspLeu.... 702
394 GCGACCGCTCGCTTCAGCAAAATTCCTGCGTGCATGCGCGAGCGCTGC 443
      |||::: |||
703 .....ProLeuGlnArgHisGlnGlySerArgGlnHisArgGlnAspArg 717
444 CATCTTCGCTCAATGCGATGAGACCAATCCGCTGCG.....TG 481
      :::::::::::::::::::: |||
718 GlnLeu...GlnGlyAspGlyHisArgLeuGlnGlyLysArgGlnLeuG 733
482 CGACCTACGCGTCACTTATCAAGAAGCGCGGAGATTTCAACGCGCG 531
      |||::: |||
733 1AlaProAlaGlyProAlaValAlaGlnAspArgArgGly..... 745
532 CTGTTGGTATGAGCGCTTTCAGCGACGCAAAATTCATGTTTGAAGC 581
      :::::::::::::::::::: |||
746 .....LysLeuArgAspArgSerGlnAlaPro..... 754
582 AGCTGGCGAGACGTGCGCTGAAATGCTGCCAACAATCGAACAACATG 631
754 ..... 754
632 AATGCGCGCGCGCATCTCGCGTTTGAGTGCACGACATTCATTTC 681
      |||::: |||
755 ...GlyArgAlaAlaArgAsnArgArgHisLeuSerAlaHisSer...A 769
682 ATGAGACCGCGTGGCGCGAATAAACCTGTGGACCATTCATTAACAAGA 731
      :::::::::::::::::::: |||
769 ArgArgAlaLeuGlnGlyProGlnGlyArgAspAlaGlyCnle..... 783
732 TGTAAATTACCATTTGCGCGTGTGTTGCAC..... 761
784 .....LeuTyrLeuProAspLeuSerArgAspArgAlaLeuArgGlnArg 798
762 ...AGCGCG...TGTGAACCGGAGCGGTGATTCGCTAGGTGTTCT 804
      |||::: |||
798 GlnArgArgProGlnGlyArgHisProAlaValAlaValLeuGlnGlyLeuA 815
805 CAAGTCAACAACCGCGCTCTTGGCTACCGTTTGGGTGC..... 845
      :::::::::::::::::::: |||
815 ArgArgArgGlnGlnLeuProAlaAspGlnProHisGlyCysGlyGlyThr 831
846 .....GAAAGTATCGCAAAATTACTCGCGCGCAATTGG 877
832 ArgArgAspGlyThrPheGlnAspPheArgAlaTyrArgGlnValAl 848
      :::::::::::::::::::: |||
878 TTGACACAGCAACCGCGTGAATTCGCGTTCGATTAAGAA..... 917
      :::::::::::::::::::: |||
848 a.....ArgLeuProArgAsnLeuPro...GlyValGlnArgSerValH 862
918 .....CGGCGGATTCACAGAGCGCGGACGATTAATTGGGACGTA 959
      :::::::::::::::::::: |||
862 1sPProHisArgArgArgHisGlnArgGlnGlyArg.....GlyArgLeu 876
960 CCACAAATCAGATTTCGTTATCGAAGAAGCGCGACAGCAAGAGCTGTGC 1009
      :::::::::::::::::::: |||
877 GlnArgHisAspArgArgHisProArgGlySerGln..... 888
1010 GCTGGGTGGCGCGGACGCGCAAAATACTCCATCAC..... 1046
      :::::::::::::::::::: |||

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889 .....AlaAspProHisGlyArgAlaLeuHisArgLeuProLeuPheG 903
1047 .....GCGTACAAACCCCTCGGCATTTCGTGA 1073
903 GlnGlnLeuProArgGlnAspAlaAlaLeuGlyPheArgValHisProArg 919
1074 AAACAAACTCTTCAAGTTCAACACACACCGCTCAACGCGCGACCGCGCA 1123
      :::::::::::::::::::: |||
920 ArgThrArgLeuSerValAsp.....GlnAlaArgArgAspArgHis 933
1124 TGTGCGGATTTGATCTTACAGCGCGTGAATGCCCTTGATTCCTGCGC 1173
      :::::::::::::::::::: |||
933 GlnAlaAspGlnGlyLeuArg.....GlnValG 943
1174 ACCCTGCTTTGGCGGATTTAATGTCGGGATACGACGCGCGCGCGC 1223
      :::::::::::::::::::: |||
943 1nPro.....GlnArgGlnArgArgHisArgArgGly 953
1224 ATGGCGTTCCTGGAATTGGA.....CGAAGAAGACCTGCTT 1261
      :::::::::::::::::::: |||
954 SerProGlnLeuGlnArgAlaValAlaHisValAlaArgHisProArgSe 970
1262 TGTGCGAG.....CTTCGCTGCGCG 1281
970 rProGlnAspAlaLysAsnProPheGlyArgTyrArgLeuAsnLeuProG 987
1282 GCGAAATACGAATACGCGCGCTGTTCGCAAGTCTGGAAC 1325
      :::::::::::::::::::: |||
987 1Y.....GlnAlaAlaGlyAsn 992
seq_name: /SIDSI/gcgdata/geneseq/geneseq_emb1/AA2001.DAT:ABB70137

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seq_documentation block:

ID ABB70137 standard; Protein; 1908 AA.

AC ABB70137;

DT 26-MAR-2002 (first entry)

DE Drosophila melanogaster polypeptide SEQ ID NO 37203.

KW Drosophila; developmental biology; cell signalling; insecticide;

KW pharmaceutical.

OS Drosophila melanogaster.

PN W0200171042-A2.

PD 27-SEP-2001.

PF 23-MAR-2001; 2001WO-US09231.

PR 23-MAR-2000; 2000US-191637P.

PR 11-JUL-2000; 2000US-0614150.

PA (PEKE) PE CORP NY.

PI Venter JC, Adams M, Li PWD, Myers EW;

DR WPI; 2001-656860/75.

PT N-PSDB; ABL14240.

PT New isolated nucleic acid detection reagent for detecting 1000 or more

PS interactions -

PS Disclosure; SEQ ID NO 37203; 21pp + Sequence Listing; English.

CC The invention relates to an isolated nucleic acid detection reagent capable of detecting 1000 or more genes from Drosophila. The invention is useful in developmental biology and in elucidating cell signalling and cell-cell interactions in higher eukaryotes for the development of insecticides, therapeutics and pharmaceutical drugs. The invention

CC discloses genomic DNA sequences (AB16176-AB130511), expressed DNA
CC sequences (AB101840-AB16175) and the encoded proteins
CC (AB57737-AB572072).
CC The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX

Sequence 1908 AA:

Alignment_scores:

Quality: 105.50 Length: 557
Ratio: 0.498 Gaps: 31
Percent Similarity: 38.061 Percent Identity: 21.544

Alignment_block:

US-09-303-518D-125 x ABB70137 ..

Align seg 1/1 to: ABB70137 from: 1 to: 1908

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859 CysProvalGlnArgGlnArgSerProserCysLysArgThrAsnPr 875
76 ATTACCGAAGTCGCGCTGCTGGCGAGAAATATGCGCGGTATCGCCCTC 125
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
875 GlnAsnArgSerArg.....LysGlnProProLecysSerArg 889
126 GATCAAGATCAAGAGGCGATGCGGCAAAAAAGCCCAAGTCG..... 169
:: :||| :||| :||| :||| :||| :||| :||| :||| :|||
889 LurThleuMeGlnLysProValLysSerTyArgPheSerCysProGlu 905
170 .....TGTTGAAGACAAAAAAGATCCGCGCGTGTGTTACTGCG 210
||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
906 AsnGlyLysCysAlaSerCysArgLysLeuLysProLysCysHisLysAl 922
211 CCGGCTTAGCGCAAAATCGCGGATTCACCGTGGCGAAAGCCGCTACT 260
||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
922 alaValaGlnLysLys..... 927
261 TCAGTCAGTCGATGATGCGCTTGAAGCAAGCAAAATGAGTTTAC 310
927 ..... 927
311 GCTACGACCTGAAGCGCTGCAAACTTAAGCGCGAAGATGCGCGC 360
:|||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
928 .....LysThrSerSerAlaLysLysCysAlaAl 937
361 AACCTGATCCAAAT.....CGGTTTGTGAGCTGCGCTGCGCACCCG 401
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
937 aSerValaLagLInLysCysCysProMetCysGlyLeu..... 949
402 TCCGTTGAGCAAAATCCCTGCGCTGATGCGGAGCCGTTGCG...CCATC 448
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
950 .....LeuProMetMetProAsnLieserLleProAsp 960
449 TCGTCATG.....CGATGACACC 468
:: :||| :||| :||| :||| :||| :||| :||| :||| :|||
961 ValGlySerGlyAsnArgLleLetyLysThrLeuGlyArgCysArgPr 977
469 AATCCGCTGCGCGCCCTACGG..... 493
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
977 o...ArgLysLeuProLysThrArgLysGluLeuThrLecCysCysLysL 993
493 ..... 493
993 ysArgCysGlnGlyArgAsnSerHisTyMetThrGlyThrValAla 1009
494 ...TCATTATCAAGAAG..... 508
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1010 ThrSerLeuLysLysArgAlaLysAlaAlaMetThrGlyValAsnProVa 1026
509 ....CCGCCGAGATTTCAAACCGCGCTGTGTTGATTCAGCCGTTTGA 554
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||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1026 lAlaProProArgValArgLInSer.....P 1035
555 CG.....ACGCAAAATCCATGTTTGTAAAGCAGCTGGCGCAG 592
|| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1035 ropHeProProLeuAsnMetProSerHisProValSerGln..... 1048
593 ACGTGGCGCTGAAATGCTGCCAACATGCACATGATTTGGCGGC 642
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1049 .....ProthGly...HisValArgSerAlase 1057
643 CCGCATCTCGCGGTTTGAAGTGGACGACATTCATTTATGAGCGGCT 692
||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1057 rArg.....AlaGlyHisSerTyLeuAspGlnAspArg 1069
693 CGGCG.....CGATAAACCGCTGTGACCATCA 721
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1069 eSerProSerThrSerAlaGlnProArgGlnArgProAlaAlaProPro 1085
722 ATTATCAAGATTAATTACCATTTGCCGCTTTGTTTCAACAG..... 764
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1086 ProPheLysMetArgSerProSerAlaAsnValaGlnLysSerGluL 1102
765 .....CCGCTGAACACCGGAGCGGTGAT 788
1102 ysArgGluGlyLysTyArgLInHisProSerGlnLInArgThrArgThr 1118
789 TGCCCT.....AGTGTGTTCTCAAGTCAACAAACCGCCCTTTC 829
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1119 MetGlnProValLleGluLInHisSerThrProLysGluGlnAlaThrAl 1135
830 GTACGCTTTGGGTGGCAAGTATGCAAAATTAATCTGCGGCAATTTGGT 879
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1135 aTyArg..... 1137
880 GACACAGACACCGCGTATTCGCGTGTGATTAAGCGCGCATTAAC 929
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1138 .....SerLInProGluProGluProSerAsnArgSerGlnArgLInPro 1152
930 ACNAGCGCGCACAGATTATTGGACGCTACACAAATCAGATTTCGGTTA 979
||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1153 ThrAsnArgSerGln.....ThrGluProProAlaSer.....Se 1164
980 TCGAAGACGCGCGCAG..... 995
1164 rArgSerArgProGluLInProAspLeuProThSerLeuLleAlaProA 1181
996 .....CAAGAGCTGTGCGCTGCGTGGCGCCGCA 1025
1181 lAserMetAspAlaGluLInLethrGluProValArgArgGlyAlaSerTy 1197
1026 GCGGCAAAATCTCCATCAGCGCTACAAACCTCGCGCATTTCCGAAAA 1075
||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1198 LeuGlyThrGluLeuAsn.....ProPheTyArgG1 1208
1076 ACAACCTTCAAGTTCAACACAGCGCTCAACGCGCGGCGCCCATG 1125
:: :||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1208 ySerPheLeuArgValArgHisSerArgAspSerProGlnSerArgL 1224
1126 GTGCGGATTTGTTACTACGAGCGCGTATGCGCTGATATCTGCCAC 1175
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1225 SerProLInHisGlnArgGlyLInLieserGluValProArgProProth 1241
1176 CTTGCTTTTCCGCGATTAATCGTGGCGGATACC.....GACAGCGCGC 1219
||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1241 rValProGluMetGlyLeuProValLysTyThrArgLeuSerGlnLeuG 1258
1220 AGCGATTGGTT..... 1231
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1258 lAlaLTrValrPheGlyAspProPheValLysGlyLysThrAsnThrG1 1274
1232 .GCTTGAATTGG.....ACGAGAAGACCTCGCTTTGTG 1265
::| :||| :||| :||| :||| :||| :||| :||| :||| :|||
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1274 yhrhrpsertrpafglniilephgilyarglysylslysserprotht 1291
1266 ca.....gcttgcgctccggcgcaatfagcaatfagccgcc 1303
1291 hrlyasprysprohalyshrhetaapalaasnaatgavmetgylgly 1307
1304 tcttgcgcgaatgctc 1318
1308 asmcysproilecys 1312

seq_name: /SIDS1/gcgdata/geneseq/geneseqp-emb1/AA2001.DAT:ABG00972

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seq_documentation_block:
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ID ABG00972 standard; Protein; 4274 AA

AC ABG00972;

DT 13-FEB-2002 (first entry)

DE Novel human diagnostic protein #963.

KW Human; chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder.

OS Homo sapiens

PN WO200175067-A2

PD 11-OCT-2001.

PF 30-MAR-2001; 2001WO-US08631.

PR 31-MAR-2000; 2000US-0540217; 00 MAR 2000 0540167

[illegible]XX
DT DE FT C

XX 2001-630363 /73

DR N-PSDB; AAS65159.

PT	New Isolated poly	diagnostics for
PT		

PT responsible for genetic disorders or other traits and to assess
PT biodiversity -
vv

PS Claim 20; SEQ ID No 31331; 103pp; English
vv

The invention relates to isolated polynucleotide (I) and polypeptide (II) sequences. (I) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chromosome and gene mapping, and in recombinant production of (II). The polynucleotides are also used in diagnostics as expressed sequence tags for identifying expressed genes. (I) is useful in gene therapy techniques to restore normal activity of (II) or to treat disease states involving (II). (II) is useful for generating antibodies against it, detecting or quantitating a polypeptide in tissue, as molecular weight markers and as a food supplement. (II) and its binding partners are useful in medical imaging of sites expressing (II). (I) and (II) are useful for treating disorders involving aberrant protein expression or biological activity. The polypeptide and polynucleotide sequences have applications in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits to assess biodiversity and to produce other types of data and products dependent on DNA and amino acid sequences. ABC00010-ABG30377 represent novel human diagnostic amino acid sequences of the invention.

Note: The sequence data for this patent did not appear in the printed specification, but was obtained in electronic format directly from WIPO at [ftp.wipo.int/pub/published_pct/sequences](http://wipo.int/pub/published_pct/sequences).

Sequence 4274 AA;

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alignment_scores:
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Quality:	105.50	Length:	454
Position:	0.533	Cans:	17

Ratio:	0.533	Gaps:	1
Percent Similarity:	43.612	Percent Identity:	21.366

Percent similarity: 40.012 Percent identity: 21.000

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alignment_block:
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US-09-303-518D-125/rev x ABG00972

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1511 ethr...ValProGlyProAlaLysSerGlyPheThrSerLeuSerS 1527
1243 CCNAATTCACAGCAACCAATGCTCGCCGCTG 1212
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1212 1212
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1560 rValLysSerIleSerAspAlaSerProIleArgSerLeuArgThr 1577
1177 GGGGGCCAGGATATCCACAGGGCATCCAGGCTCGTAA 1140
1577 etherSerProIleLysThrValValSerGlnSerProTyrAsnIleGln 1593
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1594 ValSerSerGlyThrLeuAlaArgAlaProAlaValThrGlnAlaThrPr 1610
1089 CTTGAGACGTTGTTTTCACAGGAATGCGCGAGGTTGTACCGCTGATNG 1040
1610 OlenuLysGlyLeuAlaSerSnrT 1619
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1619 hrPheSerSerArgThrSerProValThrThrAlaGlySerLeuLeuGlu 1633
989 CTTTCTTCGATACGGAATCTGATTGTGTAGCGTCCCAATATATCTGT 940
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939 CGCGCCTGTGTAATCGCGCGCTTCATACCGAACCGGAATCAGCGGT 890
1641MetThrProAlaSerProLysSerIleAsnMetT 1654
889 TGTGTGTGTCAACCAATTCGCCCGCAGTAATTTGGATCACTTTCGACCC 840
1654 YrSerSerSerLeuProPheLysSerIleIleThrSerAla...AlaPro 1666
839 AAAACGATCGCAGAGGCGCGGTTTGTGTACTGAGAACAACCTTAGGCG 790
1670LeuIleSerSerProLeuLysSe 1677
789 AATCAGCGGCTCGGTTCACAGCGCCGTTCACAAACAAAGGCCCAATG 740
1677 rValAlaSerProValLysSerArgValAspV 1688
739 TAATTAACATCTGTAATGATGATGCACACGCGTTTATTCGCGCG... 693
1688 alIleSerSerAlaLysIleThrMetAlaSerSerLeuSerSerProVal 1700
692ACGCGCTCGATGAA 679

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1705 LysGlnMetProGlyHisAlaGluValAlaLeuValAsnGlySerIleSe 1721
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649 GATGC.....GGGCGCGCGAAT 633
1734 LysLysAlaThrAlaThrLeuGlnGluLysIleSerSerAlaThrAsn 1750
632 TCATGCTTCGATGTGACGATTCACAGCGC..... 597
1751 SerValSerSerValValSerAlaAlaThrAspThrValGluLysValPh 1767
596 .....ACGT 593
1767 SerThrThrThrAlaMetProPheSerProLeuArgSerTyrValIleSerA 1784
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1784 laAlaProSerAlaPheGlnSerLeuArgThrProSerAlaSerAlaLeu 1800
542 AATACCAACAGCGCGCTTGAATCCCGCGGCTTCTTGATATGAC 493
1801 TyrThr.....SerLeuLysSerIleSerAlaTh 1811
492 CGTAGGTCGCGACGACGCGGATGTGTCATC..... 459
1811 rThrSerSerValThrSerSerIleIleThrValProValLysSerValY 1828
458 .....GCATTGACGAAGATG.....GCCACGGC 435
1828 alaAsnValLeuProGlnProAlaLeuLysLysLeuProAspSerAsnSer 1844
434 TCGGCATCGACGCGAGAAATTTGCTGACGAGCGGTGCGACGCGAGT 385
1845 PheThrLysSerAlaAlaAlaLeuLeuSerProIleLysThrLeuThr 1861
384 CCACAAACCGGATGTGATCAGCTTGGCGGCACTTTCGCCGCTTAAGT 335
1861 rGluThrHisProGlnProHisPheSerArgThrSerProValLys 1878
334 TTGCC.....AGCGTTTCAGGTCGTAGGTTCAACACTGATTTGCTG 291
1878 erSerLeuPheLeuAlaProSerAlaLeuLysLeuSerThrProSerSer 1894
290 TTGCCTTCAACG 279
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seq_name: /sids1/gcgdata/geneseq/geneseqp-emb1/AA2001.DAT:ABG07375
seq_documentation_block:
ID ABG07375 standard; Protein; 4386 AA.
XX
AC ABG07375;
XX
XX
DT 13-FEB-2002 (first entry)
XX
DE Novel human diagnostic protein #7366.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.

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PR 23-AUG-2000; 2000US-0649167.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Drmanac RT, Liu C, Tang YT;
XX
XX WPI: 2001-639362/73.
XX N-PSDB; AAS71562.
XX
XX New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits and to assess
XX biodiversity
XX
XX Claim 20; SEQ ID No 37734; 103bp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX (II). (II) is useful for generating antibodies against it, detecting or
XX quantitating a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. ABG00010-ABG30377 represent novel human
XX diagnostic amino acid sequences of the invention.
XX Note: The sequence data for this patent did not appear in the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pcl_sequences.
XX
XX Sequence 4386 AA:

alignment_scores:
Quality: 105.50 Length: 454
Ratio: 0.533 Gaps: 17
Percent Similarity: 43.612 Percent Identity: 21.366

alignment_block:
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Align seg 1/1 to: ABG07375 from: 1 to: 4386

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1504 LysProPhePheSerThrArgProTyrGlnSerThrThrAlaProIle 1520
1293 TTCGTAATTTGCCGCGGACGAGCAAGCTGCACAAAGCAGGTCTTCTGCT 1244
::: ||||| ::||| |||||
1520 eThr...ValProGlyProAlaLysSerGlyPheThrSerLeuSerSers 1536
1243 CCAATTCACAAACCAATGCTTCGCGCTG..... 1212
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1536 erSerSerAsnThrProSerAlaSerProLeuLysSerIleTyrSerVal 1552
1212 ..... 1212
1553 SerThrProSerProIleLysSerThrLeuGluValaSerThrThrSer 1569
1211 .....TCGTAATCGCGAGCAATTAATCGCGCAAAAGCA 1178
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1569 rValLysSerIleSerAspAlaSerProIleArgSerLeuArgThrW 1586
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1603 ValSerSerGlyThrLeuAlaArgAlaProAlaValThrGluAlaThr 1619
1089 CTTGACAGATTGTTGTTTTCAGAAATGCGGAGGTTGTACGCGGTATGG 1040
    |||||:||||| :|||:
1619 OleuLysGlyLeuAlaSerAsnSer.....T 1628
1039 AGATTGTGCGGCGTGCAGCAACCGCAACGACCTTTGCTGCGG 990
    :||| :|| :|| :|| :|||:|||||:
1628 hrPheSerSerArgThrSerProValThrThrAlaGlySerLeuLeuGlu 1644
989 CCTTCTCGATTACGGAATCTGATTGTGTAGCGTCCCAATATATCGTG 940
    |||||:||||| :|||:
1645 ArgSerSerIleThr..... 1649
939 CGCGCCTTGTTGTAATCGCGCTTCAATACCAACGGAATCAGCGGT 890
    :||| :|| :|| :|| :|||:
1650 .....MetThrProAlaSerProLysSerAsnIleAsnMet 1663
889 TGTGTGTACCAATTCGCCGACATTTGCCATCTTTCGACCC 840
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1663 yrSerSerSerLeuProPheLysSerIleIleThrSerAla..AlaPro 1678
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1714 LysGlnMetProGlyHisAlaGluValAlaLeuValAsnGlySerIleSe 1730
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632 TCATGTGTTCGATGTGGCAGCATTTTCAGACGC..... 597
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